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Ureteral stenosis secondary to common iliac aneurysm: a case report and review of the literature in Japan

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URETERAL STENOSIS SECONDARY TO COMMON ILIAC ANEURYSM: A CASE REPORT AND REVIEW OF THE LITERATURE IN JAPAN

Kazuo Gohji, Hideki Uehara, Shinsuke Takagi, Soichi Aракава, Osamu Matsumoto and Sadao Kamidono

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A case of right ureteral obstruction secondary to right common iliac aneurysm presenting as right hydronephrosis is reported and the literature is reviewed. A 66-year-old man was admitted to our hospital with lower abdominal pain and was treated by bypass operation with artificial vessel and uretero-ureterostomy of right ureter under the diagnosis of right ureteral stenosis secondary to right iliac aneurysm. Histological examination showed arteriosclerotic aneurysm and a fibrillary inflammatory change in peri-ureteral tissue without any hemosiderin-laden macrophages. Intravenous pyelography showed improvement of right hydronephrosis 45 days after operation.

To date 11 cases including our case have been reported in Japan, 10 in males and the other in a female. Initial symptoms were either lumbago or an abdominal mass with palpitation in many cases. The treatment was by resection of aneurysm with bypass grafting or ureterolysis.

Key words: Ureteral stenosis, Aneurysm

INTRODUCTION

The iliac aneurysm is commonly complicated with abdominal aortic aneurysm, but solitary iliac aneurysm which tends to be ruptured is relatively rare. The symptoms of iliac aneurysm were severe abdominal pain and shock following its rupture. Without rupture, it induced obstruction to adjacent organs. Ureteral obstruction, which is induced by iliac aneurysm, is rare. To date, only ten cases have been reported in the literature in Japan. We describe a 66-year-old man with right ureteral stenosis induced by right common iliac aneurysm. This is the 11th case documented in Japan.

CASE REPORT

A 66-year-old man was admitted to our hospital with lower abdominal pain. There was no evidence of hematuria, proteinuria, weight loss or fever up. Physical examinations and laboratory data were within the normal limits. Drip infusion pyelography did not show any abnormal calcification in urinary tract and demonstrated right hydronephrosis (Fig. 1). Retrograde pyelography showed a right ureteral stenosis at the level of the fifth lumbar vertebra (Fig. 2). Abdominal computerized tomography showed a mass of peri-vascular area, which was enhanced as well as aorta.

Fig. 1. Drip infusion pyelography showed moderately right hydronephrosis.
Digital subtraction angiography showed an aneurysm of right common iliac artery (Fig. 4). Replacement of the aneurysm by artificial vessel between aorta and right external iliac artery and resection of narrow segment and uretero-ureterostomy of right ureter were performed under the diagnosis of right ureteral stenosis induced by right common iliac aneurysm (Fig. 5, A, B). Intravenous pyelogram showed improvement of right hydronephrosis 45 days after the operation (Fig. 6). Histological examination

Fig. 2. Retrograde pyelography showed a right ureteral stenosis at 5th lumbar vertebra.

Fig. 3. Abdominal computerized tomography showed a mass of peri-vascular area.

Fig. 4. Digital subtraction angiography showed a right common iliac aneurysm.

Fig. 5. Pre- and post-operative status of our case (schema). A: pre-operative schema. B: post-operative schema. Replacement of aneurysm by artificial vessel, 10 mm. Goatex, between aorta and right external iliac artery and resection of narrow segment and uretero-ureterostomy of right ureter were performed. Right internal iliac artery was ligated.
Table 1. Ureteral obstruction and stenosis secondary with common or internal iliac aneurysm reported in Japan

<table>
<thead>
<tr>
<th>Case No.</th>
<th>Year</th>
<th>Authors</th>
<th>Age of patient (year)</th>
<th>Sex of patient</th>
<th>Site of Hydronephrosis</th>
<th>Site of Aneurysm</th>
<th>Initial Symptom</th>
<th>Treatment</th>
<th>Survival after diagnosis</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>1975</td>
<td>Hiekata et al.</td>
<td>22</td>
<td>M&lt;sup&gt;a&lt;/sup&gt;</td>
<td>rt&lt;sup&gt;c&lt;/sup&gt;</td>
<td>rt-IIIA&lt;sup&gt;b&lt;/sup&gt;</td>
<td>lower abdominal pain</td>
<td>resection of aneurysm</td>
<td>alive</td>
</tr>
<tr>
<td>2</td>
<td>1978</td>
<td>Gotoh et al.</td>
<td>51</td>
<td>M</td>
<td>lt&lt;sup&gt;d&lt;/sup&gt;</td>
<td>lt-CIA&lt;sup&gt;e&lt;/sup&gt;</td>
<td>lt-lumbago</td>
<td>resection of aneurysm</td>
<td>alive</td>
</tr>
<tr>
<td>3</td>
<td>1982</td>
<td>Hayashi et al.</td>
<td>61</td>
<td>M</td>
<td>bil&lt;sup&gt;f&lt;/sup&gt;</td>
<td>Ao&lt;sup&gt;b&lt;/sup&gt;</td>
<td>abdominal mass with palpitation</td>
<td>bypass grafting</td>
<td>alive</td>
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<tr>
<td>4</td>
<td>1982</td>
<td>Nishida et al.</td>
<td>68</td>
<td>M</td>
<td>rt</td>
<td>Ao</td>
<td>abdominal pain</td>
<td>bypass grafting</td>
<td>alive</td>
</tr>
<tr>
<td>5</td>
<td>1983</td>
<td>Yoshihara et al.</td>
<td>77</td>
<td>F&lt;sup&gt;b&lt;/sup&gt;</td>
<td>rt</td>
<td>rt-CIA</td>
<td>lumbago</td>
<td>no therapy</td>
<td>alive</td>
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<tr>
<td>6</td>
<td>1984</td>
<td>Nishimura et al.</td>
<td>69</td>
<td>M</td>
<td>lt</td>
<td>bil-CIA</td>
<td>lumbago</td>
<td>ureterolysis</td>
<td>alive</td>
</tr>
<tr>
<td>7</td>
<td>1985</td>
<td>Takahashi et al.</td>
<td>63</td>
<td>M</td>
<td>lt</td>
<td>lt-CIA</td>
<td>abdominal mass with palpitation</td>
<td>bypass grafting</td>
<td>-</td>
</tr>
<tr>
<td>8</td>
<td>1987</td>
<td>Tukamoto et al.</td>
<td>62</td>
<td>M</td>
<td>lt</td>
<td>Ao</td>
<td>abdominal pain</td>
<td>bypass grafting</td>
<td>alive</td>
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<td>9</td>
<td>1987</td>
<td>Yazawa et al.</td>
<td>78</td>
<td>M</td>
<td>rt</td>
<td>bil-CIA</td>
<td>none</td>
<td>bypass grafting</td>
<td>alive</td>
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<td>10</td>
<td>1987</td>
<td>Okabe et al.</td>
<td>73</td>
<td>M</td>
<td>bil</td>
<td>bil-CIA</td>
<td>lumbago</td>
<td>ureterolysis</td>
<td>-</td>
</tr>
<tr>
<td>11</td>
<td>1987</td>
<td>Goji et al.</td>
<td>66</td>
<td>M</td>
<td>rt</td>
<td>rt-CIA</td>
<td>lower abdominal pain</td>
<td>bypass grafting</td>
<td>alive</td>
</tr>
</tbody>
</table>

<sup>a</sup> M : male,  <sup>b</sup> F : female  <sup>c</sup> rt : right  <sup>d</sup> lt : left  <sup>e</sup> bil : bilateral  <sup>f</sup> IIA : internal iliac artery  <sup>g</sup> CIA : common iliac artery  <sup>h</sup> Ao : aorta
Fig. 6. Intravenous pyelography showed improvement of right hydronephrosis 45 days after operation.

Fig. 7. Histological findings of peri-ureteral tissue was fibrillary inflammatory change (H & E. x 40).

showed an arteriosclerotic aneurysm and fibrillary inflammatory changes in peri-ureteral tissue without any hemosiderin-laden macrophages (Fig. 7).

DISCUSSION

Ureteral obstruction or stenosis induced by common or internal iliac aneurysm is rare. Eleven cases including our case have been reported in the literature in Japan (Table 1). Among the 7 cases treated by bypass grafting, one patient underwent nephrectomy and transurethral resection of bladder tumor in addition, because ureter and bladder tumors were suspected. Only ureterolysis was performed in two cases. The other one case was not treated. In our case, resection of aneurysm, bypass operation with artificial vessel and uretero-ureterostomy of right ureter were performed. Rather than a function of the size of the aneurysm in itself, obstruction usually is caused by retroperitoneal fibrosis which encases the ureters. In formation of fibrous tissue, two different ways have thus far been described. First, microscopic perforations of the arteriosclerotic plaques allow blood to leak into the retroperitoneum resulting in an inflammatory process. Second, inflammatory changes usually seen in the wall of aneurysm may extend to the adventitia and perivascular structure. Although in our case, periaortic fibrosis and chronic inflammatory change were seen, we could not detect any hemosiderin-laden macrophages. Therefore, the etiology of our case suggested the above second process.

REFERENCES

Gohji et al.: Ureteral stenosis secondary to iliac aneurysm


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和文抄録

総腸骨動脈瘤による尿管狭窄の1例：本邦報告例の臨床病理学的検討

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松本 修 守殿 貞夫

右総腸骨動脈瘤による右尿管狭窄をきたした1例を報告するとともに本邦報告例につき検討を加えた。

患者は66歳男性で下腹部痛を主訴として当科受診した。右総腸骨動脈瘤による尿管狭窄の診断のもとに人工血管を用いたバイパス手術を、尿管狭窄部を切除後、尿管端々吻合術を施行した。組織学的に粥状動脈瘤と、尿管周囲に炎症性線維性組織をみとめたが、ヘモジデリンを食した大細胞癌を認めなかった。術後45日目のIVPで右水腎症の著明な改善が認められれた。

動脈瘤による尿管狭窄症の本邦報告例は自験例を含め11例認められる。その性別は、男性10例、女性1例と圧倒的に男性に多い。初発症状は、多くは腰痛あるいは排尿を有する腰部腫瘤を伴っていたが、治療は多くの例で、人工血管を用いたバイパス手術および尿管除去が施行されていた。

（泌尿紀要 34：1799-1803，1988）