A case of renal cell carcinoma found after skin metastasis

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Citation

泌尿器科紀要 (2004), 50(4): 239-243

Issue Date

2004-04

URL

http://hdl.handle.net/2433/113356

Type

Departmental Bulletin Paper

Textversion

publisher

Kyoto University
A CASE OF RENAL CELL CARCINOMA FOUND AFTER SKIN METASTASIS

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A case of renal cell carcinoma, found after skin metastasis is presented. A 79-year-old man visited Osaka JR hospital, complaining of a painless nodular mass on his right chest. The mass was resected and histopathological examination revealed a clear cell carcinoma (alveolar type, G1) with no involvement of the mammary gland. Abdominal ultrasound and magnetic resonance imaging revealed a heterogenous lower pole mass in the right kidney. Ultrasound-guided needle biopsy of the right renal mass was performed for histopathological diagnosis, which was clear cell carcinoma (alveolar type, G1). At that time, multiple metastases appeared in bilateral lung fields. The patient is currently receiving interferon-α therapy, without surgical treatment.

Key words: Renal cell carcinoma, Skin metastasis

INTRODUCTION

Renal cell carcinoma (RCC) is a significant lesion accounting for most malignant renal parenchymal neoplasms in adults and has been reported at all ages with a peak in the sixth decade. Hematoma, abdominal mass, costovertebral angle pain, weight loss, weakness and fever are the most common symptoms of the carcinoma. Most carcinomas are hypervascular tumors that tend to spread either by direct invasion through the renal capsule into the perinephric fat and adjacent visceral structures or by direct extension into the renal vein. The biologic behavior of RCC is characteristically variable and the prognosis is unpredictable.

Clear cell carcinoma is the most common histological subtype, found in approximately three-quarters of all RCC. Some renal cell carcinoma may metastasize in the early phase, but skin metastasis, as the first detected lesion of this carcinoma type is still rare.

In this paper, a case of clear cell carcinoma in the right kidney found after skin metastasis detection is presented.

CASE REPORT

A 79-year-old man visited Osaka JR Hospital, complaining of a painless nodular mass in his right chest. The mass was about 1 cm in diameter, smooth surface and was not fixed to skin. He had suffered from diabetes mellitus and diabetic nephropathy for many years, and serum blood biochemistry showed hyperglycemia and high serum levels of creatinine (3.1 mg/dL). He was admitted to Osaka JR Hospital and chest computerized tomography (CT) revealed a heterogeneous solitary mass in subcutaneous tissue on the right chest wall (Fig. 1). The mass was resected for histopathological diagnosis. The resected mass was surrounded by adipose tissue (Fig. 2), and histopathological examination revealed a

Fig. 1. Chest CT revealed a heterogenous solitary mass (arrow) in subcutaneous tissue on right chest wall.
clear cell carcinoma (alveolar type, G1); the tumor was encapsulated by fibrous and adipose tissues that were neither mammary glands nor lymph nodes (Fig. 2). The tumor was considered to be present in chest cutaneous tissue and suspected to be a metastatic lesion; therefore he was referred to the Department of Urology for further examination. Abdominal ultrasound and magnetic resonance imaging (MRI) revealed a heterogeneous lower pole mass (20×20 mm) in the right kidney (Fig. 4). Because of hemodialysis therapy for treatment of chronic renal dysfunction, the patient was introduced to Ohno Memorial Hospital, and ultrasound-guided needle biopsy of right renal mass was performed. Histopathological diagnosis was clear cell carcinoma of the right kidney (alveolar type, G1) (Fig. 5). At the same time, chest computerized tomography revealed multiple metastases in bilateral lung fields. The patient is currently receiving interferon-α (IFN-α) therapy.

Discussion

RCC is the fourth most common tumor to metastasize and approximately one-third (40–50%) of the patients have metastatic disease at the time of diagnosis. Of these patients, 1.6–3.6% will have solitary metastasis. The most common sites are lungs (50%), bone (33%) and contralateral kidney (26.5%). Regional lymph nodes, liver, adrenals and brain are also frequently involved; but, cutaneous metastasis from renal cell carcinoma is believed to be rare. The metastases may arrive through the lymphatic or the blood stream.

Brownstein et al. reviewed the 724 cases of metastatic skin carcinoma in the world and found that RCC was the fifth major cause of it in male humans, and Kishida et al. also reviewed the 62 cases of skin metastasis of RCC reported in Japan. Twenty two cases of RCC found after skin metastasis as in our case have been reported in Japan (Table 1). The age of these patients ranged between 55 and 70 years.
Table 1. Case of RCC found after skin metastasis in Japan

<table>
<thead>
<tr>
<th>Patient No.</th>
<th>Reporter</th>
<th>Year of reported</th>
<th>Age</th>
<th>Sex</th>
<th>Location of skin metastasis</th>
<th>No. of skin metastasis</th>
<th>Size</th>
<th>Location of other metastasis</th>
<th>Primary lesion</th>
<th>Treatment of metastasis</th>
<th>Survival after detection of metastasis</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Imai et al.</td>
<td>1958</td>
<td>57</td>
<td>Male</td>
<td>Right upper limb</td>
<td>1</td>
<td>Hen's egg size</td>
<td>Lung ?</td>
<td>Left kidney</td>
<td>Skin biopsy only</td>
<td>30 months (death)</td>
</tr>
<tr>
<td>2</td>
<td>Morita et al.</td>
<td>1963</td>
<td>63</td>
<td>Male</td>
<td>Left temple</td>
<td>1</td>
<td>25 mm</td>
<td>Lung. bone</td>
<td>Right kidney</td>
<td>Skin biopsy only</td>
<td>6 months (death)</td>
</tr>
<tr>
<td>3</td>
<td>Honma et al.</td>
<td>1963</td>
<td>55</td>
<td>Male</td>
<td>Right shoulder, chest, abdomen</td>
<td>20</td>
<td>Soy bean size</td>
<td>Not shown</td>
<td>Right kidney</td>
<td>Skin biopsy only</td>
<td>Not shown</td>
</tr>
<tr>
<td>4</td>
<td>Kirimoto et al.</td>
<td>1965</td>
<td>60</td>
<td>Male</td>
<td>Face, head</td>
<td>2</td>
<td>Not shown</td>
<td>Bone ?</td>
<td>Not shown</td>
<td>Skin biopsy only</td>
<td>5 months (death)</td>
</tr>
<tr>
<td>5</td>
<td>Saitoh et al.</td>
<td>1967</td>
<td>62</td>
<td>Male</td>
<td>Not shown</td>
<td>1</td>
<td>Not shown</td>
<td>Not shown</td>
<td>Right kidney</td>
<td>Skin biopsy only</td>
<td>Not shown</td>
</tr>
<tr>
<td>6</td>
<td>Kon et al.</td>
<td>1968</td>
<td>48</td>
<td>Male</td>
<td>Left back</td>
<td>1</td>
<td>Not shown</td>
<td>Lung ?</td>
<td>Left kidney</td>
<td>Skin biopsy, angiography</td>
<td>Death</td>
</tr>
<tr>
<td>7</td>
<td>Miyoshi et al.</td>
<td>1973</td>
<td>48</td>
<td>Male</td>
<td>Right temple</td>
<td>1</td>
<td>20 mm</td>
<td>Not shown</td>
<td>Right kidney</td>
<td>Skin biopsy, angiography</td>
<td>Death</td>
</tr>
<tr>
<td>8</td>
<td>Nohara et al.</td>
<td>1974</td>
<td>48</td>
<td>Male</td>
<td>Right temple</td>
<td>1</td>
<td>25 mm</td>
<td>Not shown</td>
<td>Right kidney</td>
<td>Skin biopsy only</td>
<td>Not shown</td>
</tr>
<tr>
<td>9</td>
<td>Oguchi et al.</td>
<td>1974</td>
<td>45</td>
<td>Male</td>
<td>Occiput</td>
<td>1</td>
<td>11 mm</td>
<td>Not shown</td>
<td>Right kidney</td>
<td>Skin biopsy, nephrectomy</td>
<td>Not shown</td>
</tr>
<tr>
<td>10</td>
<td>Masuda et al.</td>
<td>1977</td>
<td>59</td>
<td>Male</td>
<td>Left chest</td>
<td>1</td>
<td>Not shown</td>
<td>—</td>
<td>Right kidney</td>
<td>Nephrectomy</td>
<td>29 months (alive)</td>
</tr>
<tr>
<td>11</td>
<td>Tomokichi et al.</td>
<td>1980</td>
<td>41</td>
<td>Male</td>
<td>Occiput</td>
<td>1</td>
<td>Not shown</td>
<td>Bone</td>
<td>Right kidney</td>
<td>Skin biopsy, nephrectomy</td>
<td>12 months (death)</td>
</tr>
<tr>
<td>12</td>
<td>Inai et al.</td>
<td>1986</td>
<td>67</td>
<td>Male</td>
<td>Left latus</td>
<td>1</td>
<td>35 mm</td>
<td>—</td>
<td>Right kidney</td>
<td>Skin biopsy, nephrectomy, IFN</td>
<td>34 months (alive)</td>
</tr>
<tr>
<td>13</td>
<td>Inai et al.</td>
<td>1986</td>
<td>71</td>
<td>Female</td>
<td>left shoulder</td>
<td>1</td>
<td>30 mm</td>
<td>Lung</td>
<td>Right kidney</td>
<td>Skin biopsy, nephrectomy, IFN</td>
<td>14 months (death)</td>
</tr>
<tr>
<td>14</td>
<td>Fujii et al.</td>
<td>1988</td>
<td>54</td>
<td>Male</td>
<td>Face</td>
<td>1</td>
<td>Not shown</td>
<td>Not shown</td>
<td>Left kidney</td>
<td>Skin biopsy, nephrectomy, IFN</td>
<td>Not shown</td>
</tr>
<tr>
<td>15</td>
<td>Akiyama et al.</td>
<td>1988</td>
<td>71</td>
<td>Male</td>
<td>Right femur, back, axilla</td>
<td>3</td>
<td>Not shown</td>
<td>Not shown</td>
<td>Left kidney</td>
<td>Skin biopsy, nephrectomy, IFN</td>
<td>Death</td>
</tr>
<tr>
<td>16</td>
<td>Narita et al.</td>
<td>1990</td>
<td>65</td>
<td>Male</td>
<td>Trunk</td>
<td>Multiple</td>
<td>Hen's egg size</td>
<td>Lung</td>
<td>Right kidney</td>
<td>Nephrectomy</td>
<td>4 months (death)</td>
</tr>
<tr>
<td>17</td>
<td>Kawai et al.</td>
<td>1990</td>
<td>64</td>
<td>Male</td>
<td>Abdomen</td>
<td>1</td>
<td>Walnut size</td>
<td>—</td>
<td>Left kidney</td>
<td>Skin biopsy, nephrectomy, IFN</td>
<td>Not shown</td>
</tr>
<tr>
<td>18</td>
<td>Minami et al.</td>
<td>1991</td>
<td>54</td>
<td>Female</td>
<td>Left femur</td>
<td>1</td>
<td>13 mm</td>
<td>—</td>
<td>Right kidney</td>
<td>Skin biopsy, nephrectomy, IFN</td>
<td>Not shown</td>
</tr>
<tr>
<td>19</td>
<td>Gotoh et al.</td>
<td>1991</td>
<td>53</td>
<td>Male</td>
<td>Right shoulder</td>
<td>1</td>
<td>Not shown</td>
<td>Adrenal gland</td>
<td>Left kidney</td>
<td>Skin biopsy, nephrectomy, IFN</td>
<td>Not shown</td>
</tr>
<tr>
<td>20</td>
<td>Umeda et al.</td>
<td>1992</td>
<td>45</td>
<td>Female</td>
<td>Head</td>
<td>1</td>
<td>9 mm</td>
<td>Lung brain</td>
<td>Left kidney</td>
<td>Skin biopsy, nephrectomy, IFN</td>
<td>6 months (alive)</td>
</tr>
<tr>
<td>21</td>
<td>Hamamoto et al.</td>
<td>1993</td>
<td>74</td>
<td>Female</td>
<td>Left temple</td>
<td>1</td>
<td>Not shown</td>
<td>—</td>
<td>Left kidney</td>
<td>Skin biopsy, nephrectomy, IFN</td>
<td>Not shown</td>
</tr>
<tr>
<td>22</td>
<td>Inoue et al.</td>
<td>1997</td>
<td>76</td>
<td>Female</td>
<td>Left chest</td>
<td>2</td>
<td>18 mm</td>
<td>Lung</td>
<td>Left kidney</td>
<td>Skin biopsy, nephrectomy, IFN</td>
<td>36 months (alive)</td>
</tr>
<tr>
<td>23</td>
<td>Mitsuhashi et al.</td>
<td>2003</td>
<td>79</td>
<td>Male</td>
<td>Right chest</td>
<td>1</td>
<td>15 mm</td>
<td>Lung</td>
<td>Right kidney</td>
<td>IFN</td>
<td>22 months (alive)</td>
</tr>
</tbody>
</table>
from 41 to 79 years old, and the mean age was 59.2 years old (Table 1). The ratio of sex was similar to that of total RCC (Table 1). The skin metastases were solitary in about 16 cases and the scalp was the most common site of skin metastasis, perhaps because the scalp is of rich vascularity but scant subcutaneous soft tissue, followed by abdomen, limb and shoulder. All reported RCC cases first identified from skin metastasis were G1\(^8\), contrary to our expectation. It was only 5 cases without the metastasis to other organs (Table 1). The prognosis of the cases with metastasis to other organs was poor, but, surgical treatment was effective in the cases without other organ metastasis and better prognosis was achieved\(^8\).

In general, most skin metastasis of RCC shows soft, clear-margined, hemisphere-shape and painless nodules\(^8\). However some lesions were with pain and their pulsation palpable\(^8\). However, a histopathological approach may be needed to make an accurate diagnosis\(^8\).

In view of the treatment of the cases of RCC found after skin metastasis, it was said that radical surgery should be selected\(^8\).

In the present case, we could not see any histopathological findings of tumor invasion into the mammary glands and the tumor was encapsulated by fibrous tissue and surrounding adipose tissue. Therefore, it should not be classified as metastasis of the mammary gland. Ability to recognize lesions similar to the present case is important for at least two reasons: 1) they can represent the first manifestation of a primitive lesion otherwise improperly recognized; and 2) it may obviate radical surgical operations in the presence of widespread disease. This paper presents a case of clear cell carcinoma of right kidney identified after skin metastasis.

REFERENCES


(Received on June 5, 2003)
( Accepted on January 2, 2004)
皮膚転移によって発見された腎細胞癌の1例

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中江 晃，中尾 暢希

症例は79歳，男性。右胸部の無痛性腫瘍の自覚を主訴として，大阪J R病院外科外来を受診した。腫瘍を
外科的に切除したところ病理組織学的診断はclear cell carcinoma（alveolar type，G1）であり，乳頭組織
との関連性は認められなかったため，胸部皮膚組織への転移癌と考えられた。全身検索を行ったところ腹部
部超音波検査およびMRI検査において右腎下極に内部不均一な腫瘍を認めた。超音波エコーダイド下に
針生検を行い病理組織診断がclear cell carcinoma
（alveolar type，G1）であったため原発巣と考えられ
た。同時期になり両側肺野に多発性の腫瘍影が出現し，また患者からは積極的治療への同意をえられ
なかったため，現在，インターフェロン-α療法を施行されている。当症例は胸部皮膚転移を契機として腎細胞癌
が発見された比較的稀な1例であり，わが国では
23例目の症例報告となる。よって文献的考察を加えて
報告した。

（泌尿紀要 50：239-243，2004）