Renal cell carcinoma metastatic to the scalp

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Abstract

Because of the asymptomatic natural history of renal cell carcinoma (RCC), by the time a diagnosis is made, metastatic disease is present in about one third of the cases. Thus, the overall survival of patients with RCC remains poor. Ultimately up to 50% of patients with RCC will develop metastases. Metastatic lesions from RCC are usually observed in the lungs, liver or bone. Metastases to the brain or the skin from RCC are rare. Here we present a patient diagnosed with RCC, found to have no evidence of metastases at the time of nephrectomy, who presented two years later with metastases to the scalp. We review the literature of patients with this rare site of metastasis and outline the overall prognosis of this lesion compared to other site of metastases from RCC.

Introduction

Renal cell carcinoma (RCC) is a cancer that originates in the proximal tubule epithelium of the kidney. It is responsible for over 95% of kidney cancers and represents about 3% of malignant tumors in adults. RCC is twice more frequent in men than women and typically presents in the 7th decade of life.¹ Hematuria, flank pain and abdominal mass are the symptoms called the classic triad for the presentation of RCC despite the fact this triad is quite infrequent as it is found only in about 10% of patients diagnosed with RCC.² RCC also presents with other insidious non-specific symptoms such as weight loss, fever, and hypertension, as well as left sided varicocele in males. It might also present with anemia as a result of erythropoietin suppression. In rare instances (less than 5%) paraneoplastic syndromes associated with the disease are present resulting in hypercalcemia or erythrocytosis. The heterogeneity of the presentation of the disease compounded by the lack of warning signs makes the detection of RCC challenging. By the time of diagnosis, metastasis had occurred in about one third of the cases.³ While RCC is known to primarily metastasize to the lungs and the liver (Figure 1), virtually any organ can be involved including skin, muscle, thyroid or pancreas.⁴ Here we report a case of RCC metastatic to the scalp two years post nephrectomy following an R0 operation.

Case Report

The patient is a 64-year-old African American man who presented with back pain in the fall 2012. Diagnostic work up for the back pain included a computed tomography scan, which showed a 7 cm incidental mass on the right kidney. He then underwent right nephrectomy and pathology studies of the mass were consistent with a 5.3 cm RCC Furham grade 3, identified grossly as a golden yellow tumor with a hemorrhagic surface showing several white and gray areas that occupied the inferior pole. The excision margins were found to be negative and it was concluded that the tumor was confined to the kidney with no involvement of the vein, the renal sinus and the local lymphovascular structures.

Later in August 2014, the patient noted a left tonsillar mass with cervical lymphadenopathy and underwent bilateral tonsillectomy after an inconclusive biopsy. The surgical pathology report concluded that the mass was a 1.0×0.4×0.4 cm carcinoma metastasis, with similar histologic features to the previous RCC. The right tonsil was normal. The excision margins were again free of carcinoma.

Subsequent repeat positron emission tomography scans through spring 2015 demonstrated constantly progressing and increased osteoblastic activity with associated active soft tissue in multiple sites including left clavicle (medial third), left sixth rib, left T012 hemivertebra, right acetabulum and ischium, and right mid femur. These foci of activity were compatible with malignant and metastatic etiology. In the summer of 2015, the patient presented with bleeding lesions of the scalp grossly visible as three heterogeneously ulcerated masses, the largest of which was on the forehead measuring 4.4×3.3 cm (Figure 2).

Pathology and immunohistochemical analyses revealed atypical cells, positive for CAM 5.2 and negative for CK7 and CD68 consistent with metastasis from a primary RCC. Late in 2015, the patient presented with fever and altered mental status due to a bacterial infection. An incidental new pulmonary mass was found in the left lower lobe, presumably another metastatic mass of his previous RCC.

During 2015, the patient was treated with everolimus. Everolimus is a mammalian target of rapamycin (mTOR) inhibitor which had previously been shown to prolong progression-free survival relative to placebo in patients with metastatic renal cell carcinoma whose disease had progressed while on vascular endothelial growth factor-targeted therapy with (tyrosine kinase inhibitors such as sunitinib or soratinib).³ However, the patient decided to discontinue treatment late in 2015 due to relentless clinical progression of his disease as evidenced by the appearance of bleeding metastasis on the scalp and worsening symptoms such as chest pain and left leg pain. The patient declined pursuing further chemotherapy or radiation therapy and opted for hospice care.

Discussion

The estimated incidence of RCC in 2015 is slightly above 61,000 cases, resulting in a mortality of about 14,000 individuals. There are about 400,000 people living with kidney cancer in the United States. While the number of cases has slowly increased over the past two decades, the number of deaths has remained constant. The 5-year overall survival following diagnosis of RCC is 80% for stage I and plummets to 8% with distant metastases.¹ The median overall survival of stage IV RCC is 11.1 months in patients in which the primary tumor is surgically removed.⁵ Metastasis to the scalp from RCC is a rare
occurrence. To date, there have been less than 50 reports of RCC with metastasis to the skin and six reported metastases to the scalp (Table 1).\textsuperscript{7-11} RCC is one of the carcinomas known to spread hematogenously along with follicular carcinomas of the thyroid, choriocarcinoma, and hepatocellular carcinoma. RCC spreads via three routes,\textsuperscript{12} the most important being via the renal vein and vena cava to the right atrium and lung. However there is a lack of data supporting the hematogenous spread from the kidney (or pelvic organs) to the scalp, although theoretically possible via the renal veins to the plexus of Batson (with low pressure and flow inversion) to vertebral veins to emissary scalp veins. While the path of spread of RCC to the head, neck and skin is still conjecture, the fact remains that skin metastases are found in 1 to 3\% of cases and the preferred sites are the scalp and the face.\textsuperscript{13,14}

In the absence of metastasis, nephrectomy is the mainstay of treatment and offers 5-year survival rates of above 90\% in patients with stage I RCC. Patients with stage II and III see a relapse rate of about 40\%, usually within 3 years post-surgical excision.\textsuperscript{15} Adjuvant therapy has thus far failed to demonstrate any benefit in disease free survival (DFS), with clinical studies still being undertaken and results yet to be reported.

In the presence of metastasis, targeted therapy is often the first option. Targeted agents usually administered are tyrosine kinase inhibitor (such as sunitinib),\textsuperscript{16} vascular endothelial growth factor inhibitor (such as bevacizumab),\textsuperscript{17} or mTOR inhibitor (such as everolimus).\textsuperscript{3} The expanding portfolio of targeted agents, albeit with numerous side effects leads to complicated management of the disease, and the prognosis remains dismal with only an extra few months of DFS (i.e. sorafenib showed a DFS of 5.5 vs. 2.8 months for placebo with no difference in overall survival).\textsuperscript{18} Targeted therapies have provided a new window in which to explore potential therapeutic pathways for treatment improvement. However, with one third of patients being diagnosed at the metastatic stage, new diagnostic modalities need to be implemented.

Considerable effort is geared toward the identification of biomarkers for early diagno-

| Author                  | Age | Gender | Race           | Time to diagnosis | Treatment                      | Overall survival |
|-------------------------|-----|--------|----------------|-------------------|-------------------------------|-----------------|
| Livingston et al.\textsuperscript{7} | 45  | Male   |                | 1 year            |                               |                 |
| Snow et al.\textsuperscript{8}    | 69  | Female | White          | 6 years 4 months  | Excision with 4 mm margin     |                 |
| Eke et al.\textsuperscript{9}     | 42  | Female |                |                   | Lost to follow up              |                 |
| Pan et al.\textsuperscript{10}    | 63  | Male   |                | 6 week            |                               |                 |
| Barry Delongchamps et al.\textsuperscript{11} | 74  | Male   | African American| 2 years           | Nephrectomy, everolimus, hospice |                 |
| Current patient          | 64  | Male   | African American| 2 years           | Nephrectomy, everolimus, hospice |                 |
sis with more accurate prognostic value. However, to date, there is no biomarker that can be used clinically to help with early identification of RCC, or to offer insight regarding the aggressiveness of a tumor in a particular patient. As evidenced by this case, and despite the absence of metastasis at the time of nephrectomy, within less than two years, multiple metastases appeared including metastases on the scalp. It has been shown that metastases to the skin can appear anywhere between 6 months to 6 years in patients in newly diagnosed with RCC.

There is a lack of data regarding the prognosis value of scalp metastases and how to best manage a patient with metastases to the scalp. A retrospective study of 278 patients shows that resection of solitary metastases from RCC is associated with a 5-year survival rate of 35 to 50% arguing in favor of metastectomy when reasonable. The management of patients with RCC continues to be a challenge from early diagnosis to treatment of complicated metastases. The present report illustrates a patient who was diagnosed incidentally, but still escaped treatment and developed metastatic disease. The metastatic lesions presented a treatment dilemma, but at the end he elected for hospice care.

Conclusions

Here we present a case of renal cell carcinoma with metastasis to the scalp, two years after nephrectomy. This is quite unusual as metastases to the skin, in the case of RCC, are quite rare. Given the mortality associated with RCC (particularly stage IV), the elusive diagnosis of the condition, the presence of metastasis in one third of cases and the poor response to chemotherapy, the management of patients presenting with RCC must include frequent and timely follow-ups. The absence of metastases, even after surgery, doesn’t preclude the appearance of metastases in the evolution of the disease, emphasizing the need for continued close monitoring of patients diagnosed with RCC. It is therefore critical particularly in the setting of primary care prevention, to stress the importance of timely and regular follow-ups in patients with RCC after nephrectomy as the median time to recurrence varies from a few months to a couple of years.

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