Renal Cell Carcinoma with Parieto-Occipital Bone Metastasis

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Abstract
One third patients with renal cell carcinoma (RCC) have metastatic disease at presentation. Eight to 15% patients present with metastases in head and neck region. Skull bone metastasis is a very rare presentation. Herein we report a case of 56 year male RCC with parieto-occipital bone metastasis. He underwent radiotherapy for skull lesion and cytoreductive nephrectomy for right renal mass. Metastatic RCC is a diagnostic dilemma especially when there is no clue in clinical presentation. An unusual vascular osteolytic lesion in head and neck in middle-aged person should be dealt with a high index of suspicion for RCC.

Clinical Image

Renal cell carcinoma (RCC) constitutes about 3% of adult malignancies and 90-95 % of neoplasm arising from the kidney [1]. Renal cell carcinoma patients are mostly asymptomatic at time of diagnosis and are detected incidentally. Approximately one third of patients present with distant metastasis. Frequently metastasis is seen in lung, bone, adrenal, liver, and contralateral kidney [2]. Head and neck is involved in around 8-15% of metastatic renal tumor and sites include paranasal sinuses, larynx, mandible, temporal bones and thyroid gland [2,3]. There are very few reported cases of renal cell carcinoma presenting as parieto-occipital bone metastasis.

Herein we present a case of a 56 years old gentleman who presented with painless, progressively increasing right sided scalp mass for 6 months. On examination it was large around 10 X 10cm, non tender parieto-occipiatal swelling leading to right ear deformity, (Figure 1). Fine needle aspiration cytology revealed metastatic epithelioid malignancy. Magnetic resonance and computerized tomography imaging of head revealed a large lobulated expansile intensely enhancing mass lesion centered in scalp over right occipital and retromastoid region causing destruction of underlying occipital bone with intracranial and extradural extension. During evaluation for the primary lesion ultrasound abdomen revealed approximately 4.8 X 4.2cm exophytic mass lesion arising from lower pole of right kidney, (Figure 2). Patient was referred to urology from neurosurgery. Whole body positron emission with contrast enhanced computerized tomography scan (PET-CECT) done for staging of primary disease revealed a FDG avid large mass (9.1 X 6.6 X 7.7cm) with intracranial and extracranial extension with involvement of right parietal and occipital bones with midline shift, FDG avid subcarinal lymph node (1.9 X 3.6 cm) and FDG avid 3.5 X 3.7 X 4.2cm mass arising from lower pole of right kidney, (Figure 2 and 3). Findings were consistent with metastatic renal cell carcinoma. Scalp lesion was subjected to Intensity Modulated Radiotherapy in a dose of 40
Gray in 10 Fractions. He underwent right laparoscopic cytoreductive nephrectomy. His post operative period was uneventful discharged on post operative day 2. Histopathology report revealed pT1b, Fuhrman’s nuclear grade 3, clear cell carcinoma, with free margins. On immunohistochemistry it was CK 7: negative. Patient is doing well at 6 months follow up. Scalp swelling has become somewhat soft with overlying alopecia of scalp.

Renal cell carcinoma rarely metastasize to head and neck bones but in any patient with skull bone metastasis and unknown primary, high index of suspicion of RCCs should be kept. The prognosis of such patient is extremely poor with 1 year survival rate of 20% [3].

Treatment intent is usually cytoreductive, immunomodulation and palliative.

**References**