Intracranial Hemangiopericytoma Focally Recurrent to the Pelvis

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ABSTRACT

A 42-year-old male with a history of recurrent intracranial hemangiopericytoma presented to the Emergency Department with urinary retention. Ensuing work-up revealed a 10-centimeter circumscribed pelvic mass that appeared anatomically distinct from the sacrum. This metastasis was found 10 years after the initial resection of his intracranial lesion. An abdominal resection was performed, and the patient recovered uneventfully. Final pathology and staining findings were consistent with metastatic hemangiopericytoma. Hemangiopericytoma is a rare mesenchymal soft tissue neoplasm with potential for late local recurrence and metastasis. Here, we review the case and discuss the clinical relevance of narrowing the diagnosis to HPC or the growing umbrella of Solitary Fibrous Tumor (SFT). To our knowledge, this is the first report of intracranial HPC metastatic to the pelvis.

Keywords: Hemangiopericytoma; Solitary Fibrous Tumor; Intracranial; Immunohistochemical; Metastatic

1. Introduction

Sout and Murray originally described hemangiopericytoma (HPC) as a distinct entity in 1942. The medical community has long presumed that HPC originates from the pericytes of Zimmerman [1]. Therefore, HPC can occur anywhere throughout the body, but most frequently occurs in the lower extremities, pelvic fossa, and retroperitoneum [2]. Accounting for only 0.2% of intracranial neoplasms, intracranial HPC remains a rare event.

Small series and case reports have documented the ability of HPC to recur locally or metastasize after long intervals. Recurrence rates at 10 and 15 years have been documented with approximately 33% and 64% of patients, respectively. The most common sites of distant recurrence include bone and lung [3].

Here, we present an unusual case of intracranial HPC, recurring locally early on and finally metastasizing to the perirectal tissue, 10 years after initial resection.

2. Case Presentation

In 2003, this 36-year-old male, with a known history of previously resected occipital lobe intracranial hemangiopericytoma, presented with an intense headache, a dilated, fixed right pupil, and emesis. A head CT revealed a large right-sided intraparenchymal bleed (Figure 1) with resultant midline shift.

Figure 1. Large right-sided intraparenchymal bleed.
ditionally, there were abundant thin-walled vessels with distinct staghorn configuration. Immunohistochemical (IHC) staining was negative for S-100 and EMA, but was positive for CD34.

The patient’s postoperative course from his craniotomy was uneventful. He received a course of 54 Gy of radiation directed to the right occipital region later that year. In 2009, on a follow up MRI, the patient was noted to have a local intracranial recurrence and was treated with an additional 30 Gy of radiation to the right occipital lobe in the area of recurrence. He continued to receive routine follow up magnetic resonance imaging of his brain.

2.1. Metastatic Presentation

In December of 2011, at age 44, the patient presented acutely with urinary retention. Follow up CT and MRI imaging revealed an 8.5 × 10 cm pelvic retroprostatic mass with heterogeneous attenuation and mass-effect on the adjacent bladder, and rightward displacement of the rectosigmoid colon (Figures 2-4). Subsequently, the patient was taken to the operating room for laparotomy and an abdominal resection. Urologic consultants were unable to place ureteral stents, since they were unable to visualize the ureteral orifices during cystoscopy. The mass was found to be anterior and lateral to the rectum. It originated from Denonvilliers’ fascia. Due to technical constraints, a step-wise low anterior resection with end colostomy was required. Initial frozen section interpretation found a spindle-cell pathology. Since the tumor was hypervascular, the blood loss was about 1.5 liters. The patient suffered no apparent perioperative complications and was discharged to home on post-operative day seven.

Given the finding of metastatic HPC and unknown margin status, an adjuvant course of radiation therapy was pursued. The patient received a total of 37.5 Gy to his pelvis. The patient is currently doing well and scheduled for continuing follow up and imaging surveillance.

2.2. Pathologic Findings

The sacral mass was composed of grossly soft to rubbery tissue, predominantly tan-pink in color with focal necrosis. It was adherent to the serosa of the rectum. Sections reveal numerous thin-walled and branching “staghorn” vessels surrounded by small ovoid to slightly fusiform cells with somewhat indistinct cytoplasmic borders (Figure 5). There was patchy CD-34 immunoreactivity (Figure 6), but CD-99, EMA, smooth muscle actin, beta-catenin, bcl-2, TLE-1, and S-100 stains were negative. Approximately twenty mitotic figures were present per ten high-power fields, and the proliferation index was be-
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3. Discussion

In his initial report in 1942, Stout described nine tumors as belonging to a new class of neoplasm, hemangiopericytoma [4,5]. Even within this initial cohort of nine, there were four separate varieties. In 1949, Stout examined an additional 25 cases of HPC, but again noted significant variability among these tumors in their size, shape, and intercellular connective tissue [6]. Ultimately, in 1953, Stout conceded that HPC could even be viewed as a diagnosis of exclusion [4]. With routine pathology not including immunohistochemistry until the 1980’s and with HPC and solitary fibrous tumor (SFT) being virtually indistinguishable on light and electron microscopy, it is possible that previously assumed clinical characteristics of HPC have been a mischaracterization.

SFT was first described in 1870 as a neoplasm, typically of the pleura and of mesothelial origin. This was finally considered a unique entity in 1931 by Klemperer and Robin. Following recognition that SFT’s could originate anywhere in the body, the number reported increased exponentially in the 1990’s. Many tumors have progressively escaped from the broad definition of HPC over the past 10 years.

HPC and SFT are currently considered similar, if not identical neoplasms occupying ends of a morphologic spectrum [7]. The traditionally defined HPC is more cellular, with more rounded cells, and has a pericytic vascular arrangement. An SFT, by contrast, has more spindle-shaped cells and a more variable cellular arrangement, including the “patternless pattern”. Although “antler” or...
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REFERENCES


