Case Report
Metastasis of renal cell carcinoma to hemangioblastoma of the spinal cord in von Hippel-Lindau disease: report of a case and review of literature

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Received February 26, 2016; Accepted May 22, 2016; Epub August 1, 2016; Published August 15, 2016

Abstract: Tumor-to-tumor metastasis is considered to be an unusual event. Here, we describe a patient with von Hippel-Lindau (VHL) syndrome with metastasis of renal cell carcinoma (RCC) to a hemangioblastoma (HAB) of the spinal cord. Histological examination revealed cells with clear cytoplasm arranged in alveolar or tubular patterns in most fields of view. In the other area, the tumor showed abundant thin-walled vessels and clear cells. Immunohistochemically, the tumor cells were positive for vimentin and CK-8/18, which indicated a metastatic lesion of RCC clear cells. The diagnosis of RCC metastasis within an HAB might be overlooked because they have similar morphological characteristics. Detailed histological analyses and immunohistochemical stains should be helpful to determine whether HAB contain metastatic RCC, especially in patients with VHL syndrome.

Keywords: Von Hippel-Lindau (VHL) syndrome, renal cell carcinoma, hemangioblastoma, tumor-to-tumor metastasis

Introduction

Von Hippel-Lindau (VHL) disease is an autosomal dominant inherited disorder, characterized by multiple hemangioblastomas (HAB) of the central nervous system and cysts or tumors of various organs such as the kidney, adrenal gland and pancreas, which is now widely acknowledged as a hereditary neoplastic syndrome caused by germline mutations of the VHL tumor suppressor gene [1, 2]. Histologically, the presence of clear cells and highly vascularized stroma are common to HAB and renal cell carcinoma (RCC) of clear cell type, thus, metastatic lesions of RCC might be confused with HAB in patients with VHL disease [3]. Positive immunohistochemical reactions for cytokeratin and vimentin in the metastatic carcinomas facilitate differentiation between these two lesions [4, 5].

Tumor-to-tumor metastasis is considered to be an uncommon event in the central nervous system (CNS), which involves a pre-existing CNS tumor with metastatic deposits from a systemic neoplasm [6]. Theoretically, HAB is a good recipient of tumor-to-tumor metastasis because of its slow growth and abundant vascularization [7]. However, few cases of HAB complicated by metastatic tumor had been reported previously. We report a 34-year-old man with (VHL) syndrome who had a HAB in the cervical cord complicated by a metastasis from RCC.

Materials and methods

A case of metastasis of RCC to HAB of the spinal cord in VHL disease was retrieved in the department of Neurosurgery, Beijing Tiantan Hospital. The tumor sample was fixed in 10% neutral buffered formalin, routinely processed, and embedded in paraffin. Hematoxylin-eosin stained sections and immunohistochemical staining for both CK8/18 and vimentin were reviewed independently by two experienced pathologists.
DNA sequence analysis of the VHL gene was performed for the patient and his daughter. Venous blood was obtained, and genomic DNA was isolated from peripheral blood leukocytes. The VHL gene was amplified by polymerase chain reaction (PCR). The following primers were used for both PCR and sequencing: 5’-TGGAAATACAGTAACGAGTTGG-3’ and 5’-GGCTTCAGACCGTGCTAT-3’, the forward and reverse primers for exon 1, respectively; 5’-TGGGATTACAGGTGTGGG-3’ and 5’-GAAGTCAGCAAAATAAGC-3’, the forward and reverse primers for exon 2, respectively; and 5’-GGTAGTTGTTGGCAAAAGC-3’ and 5’-TCAAAGAGGAGAAAGACTTG-3’, the forward and reverse primers for exon 3, respectively. The PCR amplification protocol included 35 cycles of 94°C for 60 s, 60°C for 60 s, and 72°C for 60 s. Sanger DNA sequencing was performed. Products were sequenced using the ABI 3700 DNA Genetic Analyzer, and then assessed using DNAMAN.

Results

Case report

In October 2015, a 34-year-old man was admitted to our hospital with multiple HAB in the cerebellum, medulla oblongata and cervical cord,
who presented with a 4-month history of progressive neck pain and numbness of the upper extremities. Upon examination, the patient had slight neck stiffness and disorders of shallow sensation. His father died from a cerebral hemorrhage when he was still unborn. Between 2004 and 2009 he had undergone two surgical procedures for HAB in the cerebellum and medulla oblongata. A T1-weighted contrast-enhanced MRI disclosed two enhancing nodule in the medulla oblongata and 5th cervical cord (Figure 1A). The abdominal contrast-enhanced CT scan showed multiple cystic masses in the pancreas (Figure 1C) and cystic and solid masses in left kidney but without any clinical symptoms (Figure 1D). The patient and his daughter accepted the genetic evaluation after informed consent and both of them were confirmed as the missense mutation (c.500G>A (p.R167Q) and c.589G>C (p.D197H)) in exon 3 of the VHL gene (Figure 2). Intraoperatively, the tumor of the 5th cervical cord had a well-defined...
plane, and a gross-total resection was accomplished. Postoperative early stage MRI confirmed no residual mass lesion in cervical cord (Figure 1B). The tumor was gray brown and measured 2.0 cm × 1.5 cm × 1.5 cm. Histologically, part of the tumor exhibited the features of HAB, being discrete with a non-infiltrative growth pattern and having thin-walled blood vessels separated by vacuolated stromal cells scattered in a hyalinized background (Figure 3A, 3C). In the other part, the tumor exhibited the features of RCC of clear cell type, which consisted of glandular structures with alveolar and tubular patterns of growth and cells with round, central nuclei and abundant clear cytoplasm. Nucleoli were focally prominent in these epithelial cells, whereas the vascular components were fewer (H&E, × 200); C. HAB, showing a non-infiltrative growth pattern and having thin-walled blood vessels separated by vacuolated stromal cells scattered in a hyalinized background (H&E, × 200); D. Clear cells of the RCC strongly immunopositive for CK-8/18 (immunohistochemistry, × 100); E. Clear cells of the RCC strongly immunopositive for vimentin (immunohistochemistry, × 200).

Discussion

VHL syndrome is an autosomal dominant disorder with almost complete penetrance that is due to germline mutations in the VHL gene on chromosome 3q25-26 [8, 9]. Its incidence is estimated at approximately 1/36000 live births [10]. It is characterized by cerebellar, retinal, spinal, and rarely peripheral nerve HAB, RCC, pheochromocytomas, paragangliomas and various other malignant and benign tumors [6, 11]. Clinical diagnosis requires a family history of the disease and one of the aforementioned tumors or the presence of a CNS/retinal HAB plus one other tumor. In this case, the patient had an indefinite family history, but he presented multiple HAB in CNS and cysts or tumors of the kidney and pancreas, in addition, genetic evaluation showed the missense mutation in exon 3 of the VHL gene. Therefore, as mentioned, the diagnosis of VHL syndrome meets clinical criteria for diagnosis.

Tumor-to-tumor metastasis is considered to be an unusual event. This phenomenon can hap-
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pen in tumor-bearing patients with concurrent or metachronous second malignant tumors. In 1984, Pamphlett defined the criteria for diagnosis of tumor-to-tumor metastasis [12]. He established three criteria. 1) The metastatic nidus must be at least partially enclosed by a rim of histologically distinct primary tumor tissue. 2) The existence of the primary carcinoma must be proven. 3) The metastatic tumor must be demonstrated to be compatible with the primary carcinoma by morphological or immunohistochemical means [12]. RCC and meningioma are the most common visceral and intracranial recipients of metastasis, respectively, although the mechanism of tumor-to-tumor metastasis is poorly understood [13]. In theory, both HAB and meningioma are good recipients of RCC because they have abundant vascularization, slow growth and often remain unnoticed for a long time [13]. Furthermore, it is possible that RCC and HAB both represent attractive sites for metastasis on the basis of their common VHL mutations and the effects of the protein product of the VHL gene (pVHL) mutations on the extracellular matrix [11]. Previous work has shown that a normal fibronectin matrix acts to prevent tumor cells from spreading into the extracellular matrix and then from spreading, thus the presence of a defective matrix may provide an environment that allows metastatic tumor cells to grow successfully [11].

Metastatic RCC develops in 40% of patients with VHL disease, and it is the cause of death in one third of them [3]. Earlier detection of RCC has an important impact on survival. Since high vascularization and clear cell morphology are common to both tumor types, these characteristics sometimes cause differential diagnostic problems between HAB and metastasis of clear cell RCC. Histologically, typical RCC has alveolar or tubular patterns, water-clear cytoplasm, and an absence of vacuolated stromal cells. HAB has prominent proliferations of stromal cells, and occasionally displays well defined lobules. However, it is often difficult to see detailed cell structure because of the abundant vascular tumor tissue and the little tissue available for examination. Thus, morphology alone may not serve as a reliable diagnosis. Immunohistochemical stains can be used to support this histological distinction. Immunohistochemical staining for cytokeratin, PAX-8, EMA, CD10 and vimentin facilitates the differentiation of HAB and metastatic clear cell RCC, as HABs are negative for these epithelial markers [5]. Additionally, the MIB-1 labeling index (Ki-67 immunostain) is reported to be low in HAB (less than 5%) but moderate to high in RCC (more than 5%) [14].

Spinal HABs make up less than 2% of all spinal cord tumors and less than 7% of all central nervous system HABs [15]. Although HABs may occur sporadically, a spinal location is strongly associated with VHL disease [15]. Metastatic RCC to HAB of the spinal cord was first reported in 2001, in a patient with VHL disease. Since then, reports of 10 more cases have been published [16]. The current case is the first report that the metastasis of RCC to a spinal HAB in Chinese patients with VHL disease.

In conclusion, RCC is the most common metastatic tumor amongst those documented to metastasize to HAB and it usually occurs in the context of VHL syndrome. The diagnosis of RCC metastasis within an HAB might be overlooked because they have similar morphological characteristics. Detailed histological analyses and immunohistochemical stains should be helpful to determine whether HAB contain metastatic RCC, especially in patients with VHL syndrome.

Acknowledgements

This study was supported by National Key Technology Research and Development Program of the Ministry of Science and Technology of China (No. 2013BAI09B03) and Beijing Institute for Brain Disorders (No. BIBD-PXM2013_014226_07_000084). Written informed consents were obtained from the patients for publication of this case report and any accompanying images. A copy of the written consent is available for review by the editor-in-chief of this journal.

Disclosure of conflict of interest

None.

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References