MRI features of intracranial anaplastic hemangiopericytoma

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Abstract. Magnetic resonance imaging (MRI) features of intracranial anaplastic hemangiopericytoma (AHPC) were analyzed. The pathological examination showed that there was a great number of irregularly arranged tumor cells with nuclear atypia, and mitotic properties were commonly seen providing support for clinical staging, therapy and prognosis judgment. Eighteen cases of intracranial AHPC proved by operation and pathology were analyzed retrospectively. Both plain and enhanced MR scans were performed and the results were compared with pathology in all cases. In all 18 cases, the tumor was positioned in the cortex; in 12 cases, it was located in the frontal falx and in 3 cases, it was located in the parietal falx. In 2 cases, the tumor was located in the middle cranial fossa, and in 1 case, it was located in the cerebellar hemispheres. Thirteen of the 18 cases showed mixed hyper-iso signal intensity with cortical grey matter, and the other 5 cases were isointense in the cortical grey matter on T1-weighted images. Fifteen of the 18 cases showed heterogeneous hyper-iso signal intensity, and the other 3 cases were isointense on T2-weighted images. Fifteen of the 18 cases showed heterogeneous enhancement in contrast-enhanced T1-weighted images. Our data show that, because intracranial AHPC has specific features on MRI, it could be very useful for its clinical diagnosis.

Introduction

Hemangiopericytoma (HPC) was first reported and named by Stout and Murray in 1942 (1). It is a rare soft tissue tumor and may occur in any part of the body, but it occurs less commonly in the central nervous system (CNS). In the past, HPC was believed to originate from the meninges and was thus considered a subtype of meningioma. In the 2000 World Health Organization (WHO) classification of CNS tumors, HPC was categorized as a non-meningeal epithelial cell tumor. It originates from Zimmerman meningeal cells in interstitial capillaries, has high cell density and multi-differentiation potential, and is highly vascularized. HPCs are classified as WHO II-III grade tumors according to the 2000 classification system. In the 2007 WHO classification of CNS tumors, HPC was divided into WHO III grade anaplastic hemangiopericytoma (AHPC) and WHO II grade HPC (2-4). AHPC is more aggressive and exhibits common recurrence and extracranial metastasis. Pathological examinations revealed a great number of irregularly arranged tumor cells that commonly have nuclear atypia and mitotic properties. In this report, we present the imaging and pathological features of 18 cases of AHPC and analyze the features of AHPC that were visualized by MRI to improve the knowledge on AHPC.

Patients and methods

Clinical data. This study included 18 cases of AHPC that were confirmed by surgery and pathology at the Lanzhou University Second Hospital from July 2001 to July 2013. Twelve male and 6 female patients participated to the study, and were from 29 to 61 years of age (mean age, 44 years). This study was approved by the Ethics Committee of the First Affiliated Hospital of Chongqing Medical University. Signed written informed consents were obtained from all participants before the study. The patients agreed to the use of their samples in scientific research.

Magnetic resonance imaging (MRI) technique. MRI scans were performed with a 1.0T scanner (Magnetom Harmony; Siemens Healthineers, Erlangen, Germany). The imaging protocol included unenhanced axial and sagittal T1-weighted sequences, axial and coronal T2-weighted sequences, and contrast-enhanced T1-weighted images. Our data show that, because intracranial AHPC has specific features on MRI, it could be very useful for its clinical diagnosis.

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Pathological examination. The tumor was collected and fixed in 4% buffered formalin for 24 h. Each fixed sample was then cut into 3-4 µm slices, to make 5-6 sections, and tissue blocks were selected from representative areas. After processing and paraffin wax embedding, the sections were separately stained with hematoxylin and eosin (H&E), CD34, Ki-67, vimentin and epithelial membrane antigen (EMA). All slides were reviewed by two neuropathologists.

Results

General data. Nine patients presented with intracranial hypertension and physical signs, and 2 of those patients had two symptoms. Thirteen cases presented with headaches and/or dizziness. Three cases exhibited homonymy hearing loss. Four cases exhibited olfactory dysfunction. Psychiatric symptoms were present in 2 cases. Epilepsy was found in 10 cases, and hemi-vision loss was found in 3 cases (Table I).

MRI findings. In all 18 cases, the tumor was positioned in the cortex; in 12 cases, it was located in the frontal falx, and in 3 cases, it was located in the parietal falx. In 2 cases, the tumor was located in the middle cranial fossa, and in 1 case, it was located in the cerebellar hemispheres. The diameter of the tumors ranged from 3.1 to 6.5 cm, with an average of 4.9 cm. The tumor mass was lobulated in 12 cases, irregular-shaped in 5 cases, and oval-shaped in 1 case. In 9 cases, the tumor was connected to the dura or skull with a narrow base. On MRI, the lesions showed a mixed high-low signal in 12 cases, an iso-signal in 6 cases on plain T2WI, a mixed iso-low signal in 11 cases, an iso-mild high-low signal in 1 case, and an iso-signal in 6 cases on plain T1WI. After contrast injection, different levels of enhancement were observed in all cases. In 3 cases, skull bone damage was found, and the peripheral edemas in those cases had high signals on T2WI. For the MRI enhanced scans, a substantial part of the tumor was significantly enhanced, and the neighboring meninges were linearly enhanced (Table II) (Figs. 1-3). Preoperative imaging diagnosed HPC in 4 cases, meningioma in 9 cases, malignant meningioma in 8 cases, a metastatic tumor in 1 case, and glioma in 1 case. All 18 cases were AHPC, as confirmed by pathological analysis after surgery.

Discussion

Intracranial HPC is a rare malignancy that usually originates from the intracranial vasculature. It accounts for only 1% of all primary CNS tumors (5). Intracranial HPC was previously believed to originate from the meninges; thus, it was considered a subtype of meningioma. However, with the development of molecular genetics, it was confirmed that HPC has a completely different source from that of meningiomas: it actually originates from arachnoid cap cells, as determined by the detection of the neurofibromatosis 2 (NF2) gene in HPC tumors (6). In 1993, WHO classified HPC as different from meningioma (7). However, a review of the literature and the data collected in this study indicated that the 1993 classification of HPC did not distinguish its subtypes (8-10). The

| Presenting symptom or sign | No. of patients | Average age (years) | Males | Females |
|----------------------------|-----------------|---------------------|-------|---------|
| Intracranial hypertension and physical signs | 9 | 52 | 5 | 4 |
| Headaches and/or dizziness | 13 | 48 | 10 | 3 |
| Hearing loss | 3 | 44 | 1 | 2 |
| Olfactory dysfunction | 4 | 51 | 3 | 1 |
| Psychiatric symptoms | 2 | 60 | 0 | 2 |
| Epilepsy | 10 | 35 | 7 | 3 |
| Hemi-vision loss | 3 | 42 | 2 | 1 |

AHPC, anaplastic hemangiopericytoma.

| MRI features | No. of patients |
|--------------|-----------------|
| Position | |
| Frontal falx | 12 |
| Parietal falx | 3 |
| Middle cranial fossa | 2 |
| Cerebellar hemispheres | 1 |
| Shape | |
| Lobulated | 12 |
| Irregular in shape | 5 |
| Oval-shaped | 1 |
| Narrow base connected to dura | 9 |
| T2WI | |
| Mixed high-low signal | 12 |
| Iso-signal | 6 |
| T1WI | |
| Mixed iso-low signal | 11 |
| Iso-mild high-low signal | 1 |
| Iso-signal | 6 |
| Bony destruction | 3 |
| Peritumoral edema | 15 |

MRI, magnetic resonance imaging; AHPC, anaplastic hemangiopericytoma.
2007 WHO classification divided intracranial HPC into two separate categories: WHO grade II HPC and WHO grade III AHPC with malignant biological behaviour (11). There are differences between these subtypes in the 5-year survival rates, recurrence rates and transferability, and some studies have shown that AHPC recurs as much as 6-7 years earlier than HPC does (12).

The average age of patients with AHPC is 44 in this study. The incidence of AHPC in males is slightly higher than in females. The symptoms of headache and intracranial pressure
in different parts of the brain are very common in AHPC patients. It results from MRI findings that the average size of the lesions was 4.9 cm. The AHPC’s malignant signs, such as tumor lobulation, necrosis and cysts, were more common because they contained a greater number of irregularly arranged tumor cells in which nuclear atypia and mitotic properties were more easily found. It often co-existed with necrosis and cystic changes, so the signal of the MR plain scan was also mixed. Necrosis and cystic properties of the tumor reflected a rapid tumor growth and a relative lack of nutrition, which indicated the characteristic of high malignancy. From Akiyama et al. opinion (13), the irregular shape and ill-defined boundary reflected the rapid growth, and with invasive growth features of malignant tumors. In addition, such signs as skull destruction and peritumoral edema were commonly found in AHPC. The sign of skull destruction indicated a strong level of invasiveness. The sign of peritumoral edema reflected the amount of infiltration within the tumor, blood supply and pathological type as reported by Lee et al. (14). The AHPC has a rich blood supply because pathological H&E staining showed that there was a large number of slit-like blood vessels and an enhanced MR scan displayed that the tumor was significantly enhanced. Immunohistochemical staining results also showed the malignant tendency of AHPC.

We found that intracranial AHPC should be differentiated from malignant meningioma after literature reviewing. Malignant meningioma has some characteristic MRI features similar to AHPC (15). i) Clinical features: malignant meningioma was more common in 50-year-old females, and AHPC was more common in 45-year-old males; and ii) MRI features: malignant meningioma was generally connected to the meninges by a wide base, and AHPC was generally connected to the meninges by a narrow base. AHPC showed a more lobulated and irregular shape and cross-leaf growth, more necrosis and cysts, less skull damage; and malignant meningioma showed a less lobulated shape, less necrosis and cysts and more skull damage. Furthermore, AHPC showed more mixed signals.

In conclusion, intracranial AHPC is a rare tumor type that has a high degree of malignancy. Recurrence and metastasis are common with AHPC. The location of AHPC is similar to meningioma, but the shape of AHPC tends to be irregular. Mixed signal is more likely to be seen and MRI enhancement often shows heterogeneous enhancement in contrast-enhanced T1-weighted images. Skull destruction and peritumoral can easily be found in AHPC. Understanding the MRI features of AHPC has great significance for guiding clinical treatment and predicting prognosis.

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