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Extraaxial Cavernous Hemangioma with Hemorrhage

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Intracranial cavernomas, initially described as rare lesions [1], have been seen with increasing frequency since the advent of MR imaging [2-5]. MR features of cavernomas are highly suggestive of the diagnosis, although not pathognomonic. Cavernomas are classically intraparenchymal lesions and occult to angiography. They have a pseudotumoral growth [6] and a low potential for life-threatening hemorrhage, at least in supratentorial locations [7].

We report a case of a vascular lesion in an atypical location that was identified as a cavernous hemangioma. The lesion arose in the perimesencephalic cistern, and was accompanied by symptoms of acute onset of diplopia and meningeval signs. Extracerebral cavernous hemangiomas, as reported in the literature, have slightly different clinical, angiographic, and surgical features compared with intraparenchymal lesions, and can easily be misdiagnosed as meningioma unless MR imaging is performed.

Case Report

A 43-year-old woman without previous neurologic disease was admitted for diplopia and headaches of 3 weeks duration. A CT study performed 2 weeks after onset of symptoms disclosed a hyperdense nodular lesion of the right perimesencephalic cistern (Fig. 1A). The patient waited 1 week before seeking medical treatment.

Clinical examination on admission revealed paresis of the right superior oblique muscle. MR imaging (Signa 1.5 T, General Electric, Milwaukee, WI) showed a round, 8-mm lesion in the perimesencephalic cistern beneath the tentorium with no connection to any of the regional vessels. The lesion had a heterogeneous hyperintense signal on both short and long TR/TE sequences, and enhanced after contrast injection (Figs. 1B and 1C). Surgical removal was not attempted at the time. The patient's symptoms remained identical for 3 months, when the headaches increased acutely in intensity and frequency. A repeat MR study showed enlargement of the previously identified lesion with signal features of a "cavernomatous" lesion. The enlarged lesion now appeared intraparenchymal and bulged into the adjacent subarachnoid space (Figs. 1D-1F).

At surgery, the lesion was found to be enclosed within a thin arachnoid membrane, which was adherent to the lateral aspect of the right cerebral peduncle. The mass was completely removed, and microscopic examination revealed a hemorrhagic lesion with internal fibrous septa suggestive of cavernous hemangioma.

Discussion

According to Russell and Rubinstein [8], cavernomas are classified along with capillary telangiectases, venous an­giomas, and arteriovenous malformations as intracranial vascular hamartomas. Cavernomas classically appear as honeycomb-like dilated vascular spaces separated by endothelial-lined fibrous septa without interspersed neural tissue or elastic or smooth muscular components. The presence of intervening neural tissue distinguishes cavernomas from capillary telangiectases. According to Simard et al. [9], the presence of muscular fibers does not exclude the diagnosis of cavernomas. Thrombosis and low internal flow are the main phenomena that account for the inability to see cavernomas on standard angiographic examinations [10]. CT is disappointing in its depiction of occult cerebrovascular malformations. In a series reported by Golomb et al. [4] of 49 patients with MR characteristics of occult cerebrovascular malformations, only 29 had a positive finding on CT.

MR features are highly suggestive of cavernomas although not pathognomonic. Findings include clusters of foci of hyperintense signal on all sequences, resulting from extracellular methemoglobin corresponding to hemorrhage in different stages of evolution, which are surrounded by a hypointense rim of hemosiderin on T2-weighted images. Hemorrhagic metastases, capillary telangiectases, and thrombosed arteriovenous malformations can exhibit the same features [11, 12].

Typical locations of cavernomas include the subcortical hemispheric area and brainstem [1, 8]. Subarachnoid, subdural, and epidural cavernomas have already been described, but represent atypical locations [8]. Only five extracerebral lesions were reported by Simard et al. [9] after a review of the literature of 126 patients, one by Voigt and Yasargil [13], and none by Lobato et al. [14] in their review of 241 cases of occult cerebrovascular malformations. Most of the extraparenchymatous cavernomas have been found in the middle cranial fossa, where they may mimic meningioma [15-20].
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Cerebellopontine angle lesions have also been reported as extending into the internal auditory canal [21]. Extracerebral cavernomas, described as highly vascularized lesions with an angiographic blush, may mimic meningioma, even on CT [18, 22]. Although most of the neuropathologic textbooks make no distinction between cavernomas and cavernous hemangiomas [1, 8, 23], Lasjaunias et al. [22], relying on the reports of Harris and Jakobiec [24] on intraorbital cavernous hemangiomas, assert that the two lesions do not correspond to the same pathologic entity. Cavernomas are intraparenchymal and can grow by iterative intrallesional microhemorrhage caused by weakening of the parietal septa by thrombosis or, less commonly, by angiectasia [25]. Cavernous hemangiomas are mainly extraparenchymatous and constitute a truly tumor disease with an in vitro potential of proliferation of endothelial cells, as in liver or orbital locations [24, 26].

The combination of angiography, CT, and MR findings should allow an easy diagnosis of the lesion. Meningioma can now be excluded, since hemorrhage is an atypical feature of small meningioma [27]. The main differential diagnoses that could remain are leptomeningial hemorrhagic metastases especially from melanoma [11] and thrombosed aneurysm when the lesion is close to a major blood vessel [28].

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REFERENCES