Thoracic extraosseous, epidural, cavernous hemangioma: Case report and review of literature

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ABSTRACT

A 40-year-old male presented with mid-thoracic backache and progressive, ascending, spastic, paraparesis for one year. Magnetic resonance imaging demonstrated an extraosseous, extradural mass, without any bone invasion at the T2-T4 vertebral levels, located dorsal to the thecal sac. The spinal cord was compressed ventrally. The lesion was totally excised after a T2-T4 laminectomy. Histopathological examination revealed a cavernous hemangioma. The authors reported this case and reviewed the literature, to explain why extraosseous, extradural, cavernous hemangiomas should be considered in the differential diagnosis of extradural thoracic compressive myelopathy.

Key words: Cavernous hemangioma, epidural, magnetic resonance imaging, myelopathy, thoracic spine

Introduction

Cavernous hemangiomas (CH) in the spine are uncommon hamartomatous malformations, which occur per se in vertebral bodies, but may have an associated extradural component. Previously described nomenclatures include cavernous angiomas, hemangiomas, varicosities, venous angiomas, and angiolipomas, with cavernous vessels.[1‑5] This may be better understood in the context that histological classification can be affected by the predominant type of vascular channels. Extraosseous, extradural cavernous hemangiomas (EECH) are relatively uncommon, differ radiologically from their cranial counterparts, and may be difficult to diagnose preoperatively.[6‑22] Purely extradural causes of compressive myelopathy are by themselves rare entities. The authors discuss EECH as a part of this differential diagnosis.

Case Report

A 40-year-old male presented with a band-like feeling of constriction at the nipple level for eight months, followed by numbness to all modalities of sensation below this level. During the last month, progressive ascending spastic paraparesis had rendered him wheelchair-bound. There was no history of trauma, fever, deformity, tenderness over the spine, weight loss, or bowel or bladder dysfunction. He had no neurocutaneous markers. He had grade 3/5 medical Research Council (MRC) spastic paraparesis, exaggerated bilateral knee and ankle reflexes, and sustained ankle clonus on the left side. The lesion was totally excised after a T2-T4 laminectomy. Histopathological examination revealed a cavernous hemangioma.
and to the right. A central hyperintensity on T2-weighted imaging, within the cord substance, was also observed. Of note was the presence of a remote intraosseous T8 vertebral hemangioma. The lesion enhanced avidly and homogenously with contrast [Figure 1]. Computerized tomography (CT) of the upper dorsal spine did not reveal any bony destruction or widening of the intervertebral foramen. A radiological differential of an extradural schwannoma or meningioma was considered.

**Surgery**

The patient underwent a T2-T4 laminectomy. Intraoperatively, the mass was encountered within the epidural fat and immediately beneath the laminae. It was reddish in color, firm, friable, and highly vascular. It was located epidurally, mostly dorsal to the thecal sac, but also extended laterally to the left, along the left T2 nerve root sheath. Its blood supply was derived from the intervertebral foramina on both sides. Initially, the tumor’s dorsal surface was devascularized, with bipolar cautery, under the operating microscope. Its arterial feeders were then sequentially cauterized and interrupted, just medial to the intervertebral foramina, taking care not to injure the dorsal nerve roots. The lesion was then peeled off the surface of the dura and the adjacent nerve sheaths. The lesion was excised en bloc. The underlying dura was then opened, taking care to preserve the arachnoid, to ensure total microsurgical excision.

The patient was discharged 48 hours after surgery. His postoperative course was significant for a subjective improvement in spasticity and numbness. The histopathological examination indicated a CH [Figure 2]. On the last follow-up, 12 months after surgery, a check craniospinal MRI confirmed complete excision, with no recurrence at the operative site and no other new lesions. His power and sensations were nearly normal and he had returned to work.

**Discussion**

**Epidemiology**

Cavernous hemangiomas are ubiquitous. Within the central nervous system, they tend to occur predominantly within the cranial cavity’s supratentorial compartment. Within the spine, CH are rare. Hillman and Bynke (1991) reported an incidence of 0.22 cases/million/year. Although EECH are supposedly rarer, slightly less than 100 cases have been reported in the literature to date, since Globus and Doshay first described this entity in 1929. In this specific subgroup, a female preponderance is noted (70%). The average age at diagnosis has been reported to be around 40 years, which is the exact age of our patient. EECH are generally seen in the thoracic spine (60%), and are typically located posterolaterally and may extend into the neural foramina (19%). An extraspinal component may be evident in about 13% of all cases. The most common presentation is that of slowly progressive paraparesis (71%) and radiculopathy (10%).

**Figure 1a:** T2-weighted, mid-sagittal magnetic resonance image revealing a thoracic, hyperintense, solitary, extradural mass (black arrows) opposite the D2-4 vertebral bodies. The signal intensity differs from that of the cerebrospinal fluid, but is similar to that of a remote intraosseous hemangioma within the D8 vertebral body. Axial magnetic resonance images reveal an eccentrically placed dorsal extradural mass (asterisk), which is isointense on T1- (b) and hyperintense on T2- (c) weighted images, with left foraminal extension (black dashed arrow)

**Figure 2:** Photomicrograph showing numerous dilated vascular channels filled with blood and separated from each other by fibrous septae. These vascular channels do not communicate with each other (H and E x200)
Investigations
An MRI is the investigation of choice and demonstrates a well-circumscribed, lobulated lesion, iso- to-hypointense to the spinal cord on T1- and hyperintense on T2-weighted images, with avid homogenous post-contrast enhancement.\[6-10,14,22\] The peripheral rim of hypointensity, resulting from hemosiderin deposition, is usually seen in the intramedullary CH and not in EECH. EECH also differs from their intracranial counterparts in that they are homogeneously hyperintense on T2-weighted images, and enhance avidly with contrast. In this, they may resemble the CH arising from the cavernous sinus more. The absence of popcorn-like peripheral blood products may be explained by the fact that these may be washed away as EECH lie outside the blood-brain barrier.\[14\] It may be impossible to differentiate between extradural CH and extradural schwannomas or neurofibromas.\[5-8,10,14\] We were certainly unable to differentiate between the two preoperatively. During surgery too, our first thought was that of a vascular metastasis, as the lesion lacked a classical multilobulated purplish mulberry/blueberry appearance and was instead reddish. This was described by other surgeons too.\[12,14,18,22\] Feng et al.\[9\] had described the visual effect of contrast enhancement within these tumors as that of ‘wafting silk’, with lobules bulging into the interlaminar space on the sagittal images, especially so in the thoracic spine. They also described widening intervertebral foramina and bone erosion. Elegant as it sounds, we have seen this appearance before in schwannomas, meningiomas, and round cell tumors, and still believe that to make a preoperative diagnosis reliably, in a classical case such as ours, it is not possible by using MRI alone. However, the addition of angiography may make a difference. EECH are angiographically occult, but enhance brilliantly with contrast. Osseous CH show up on angiograms as would vascular schwannomas and meningiomas.\[23\]

Conclusions
Extrasosseous, extradural cavernous hemangiomas are relatively rare lesions within the vertebral canal. They occur most commonly in middle-aged women, in the thoracic spine, dorsal to the spinal cord, but are eccentrically placed. Intervertebral and extraforaminal extensions can occur. EECH frequently mimic the more common schwannomas clinically and radiologically. On an MRI, classical EECH signal properties differ significantly from their intracranial and intramedullary counterparts and more closely resemble the cavernomas of the cavernous sinus. Although preoperative differentiation may not be reliable using the current imaging techniques, EECH must form an important differential diagnosis of a purely extradural thoracic compressive lesion. Surgical excision in expert hands results in an excellent outcome. Adjuvant SRS may be effective in residual lesions after subtotal removal.
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How to cite this article: Sharma MS, Borkar SA, Kumar A, Sharma MC, Sharma BS, Mahapatra AK. Thoracic extraosseous, epidural, cavernous hemangioma: Case report and review of literature. J Neurosci Rural Pract 2013;4:309-12.

Source of Support: Nil. Conflict of Interest: The authors declare no potential conflict of interest.