Epidermoid Intramedullary Cyst: A Rare Case Report

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Abstract

Epidermoid cysts are tumors that occur in most parts of the body and are often treated by neurosurgeons, but intramedullary localizations of epidermoid cysts are very rare. The incidence in adults is lower than 1% and in children lower than 3%. We report a case of a 27-year-old patient presenting with thoracic spinal cord compression syndrome (T11-L2). The magnetic resonance imaging (MRI) showed an intramedullary cystic mass. It demonstrated low signal intensity on T1-weighted sequence and high signal intensity on T2-weighted sequences, with minimal peripheral enhancement following intravenous administration of gadoteric acid. After surgical removal of the tumor, the pathology confirmed the radiological diagnosis of spinal epidermoid cyst. MRI reduces the delay in diagnosis of spinal cord tumors but should be guided by clinical judgment.

Keywords: Epidermoid cyst; Spinal cord cyst; Spinal cord; Magnetic resonance imaging

Introduction

Epidermoid cysts are a frequently occurring subgroup of congenital tumors. In a series of adult patients, cranial site involvement outweighed the spinal sites by 14:1. Intraspinal epidermoid cysts represent less than 1% of all intraspinal tumors in adults [1].

In children, the incidence is higher. However, most intraspinal epidermoid cysts are subdural and extramedullary, and they are rarely found within the spinal cord. We review the literature regarding intramedullary epidermoid cysts and present the case of a child with an intramedullary epidermoid cyst in the thoracic region that was evaluated by magnetic resonance imaging (MRI).

The vast majority of spinal epidermoid cysts are localized at subdural and extramedullary spaces. Thoracic region is the favorite site of the intramedullary epidermoid cysts and the lumbar region is the next common area [2].

The differential diagnosis depends on the signal intensity of cyst content and the enhancement pattern of cyst wall. However, because of the similar imaging features, it is not easy to distinguish this pathology among other entities that are usually encountered in differential diagnosis during MRI examinations (epidermoid cyst, dermoid cyst, ventriculus terminalis and arachnoid cyst).

In conclusion, epidermoid cysts should be included in the differential diagnosis when an intramedullary cystic mass with the abovementioned MRI features is detected within the spinal canal.

Case Report

A 27-year-old woman developed progressive difficulty in walking associated with stiffness and paresthesia during 3 months. There was no history of trauma or any medical procedure on the spine. Clinical (neurological) examination revealed spastic paraparesis with the grade 1 muscle power (Medical Research Council grading scale) on both lower limbs. Sensory examination revealed loss of all sensation below T11-L1 segments. Sacral dermatomal sensations were preserved. Deep tendon reflexes decreased in both lower limbs. MRI study of the thoracic and lumbar spine was performed. An intramedullary cystic mass measuring 16 mm (transverse plane) and 66 mm (craniocaudal diameter) at the T11-L1 levels was found. The lesion was hypointense on T1-weighted sequence (Fig. 1a) and hyperintense on T2-weighted sequence. Turbo inversion recovery magnitude (TIRM) sequence, T2 fat sat and T1 fat sat (Fig. 1b-d and Fig. 2a-c) showed minimal peripheral enhancement following intravenous administration of gadoteric acid (Fig. 3). The patient underwent T11-L1 laminectomy with removal of the tumor. The slices obtained during histopathology examination showed a fibrous wall lined with stratified ceroic epithelium surrounding a cyst, consistent with an intramedullary epidermoid cyst (Fig. 4). After surgical removal of the tumor, the pathology diagnosis confirmed the MRI diagnosis of spinal epidermoid cyst. Figure 5 shows T2-weighted sagittal sequence (segments T11-L1) after surgical removal of the tumor (Fig. 5a), T1 fat sat sagittal sequence (segments T11-L1) after surgical removal of the tumor (Fig. 5b), T1 fat sat coronal sequence (segments T11-L1) (Fig. 5c), and T1 fat sat transverse sequence after administration of intravenous contrast (segments T11-L1) and after surgical removal of the tumor (Fig. 5d).

Discussion

Intramedullary epidermoid cysts with localization in the spi-
Epidermoid Intramedullary Cyst

The vast majority of epidermoid cysts are localized at subdural and extramedullary spaces. Thoracic region is the most favorite site of intramedullary epidermoid cysts and the lumbar region is the next common area of occurrence [2].

Histologically, epidermoid cysts have a fibrous wall lined with stratified squamous epithelium surrounding a cyst containing waxy squames [3]. It can be either congenital or acquired.

As for the epidemiology, the incidence of the epidermoid cysts among other intracranial tumors in most of the large series of cases is estimated to be from 0.2% to 1%; in the spinal canal, the incidence is even lower [4].

Halcrow et al reported a series of 90 intraspinal epidermoid cysts collected from the literature, with 39 being acquired and 51 being congenital. Acquired epidermoid cysts have been found years after single or multiple lumbar spinal punctures and are thought to result from iatrogenic penetration of skin fragments [5, 6]. It is generally believed that

Figure 1. (a) T1-weighted sagital sequence (T11-L1 segments show hypointense, heterogeneous signal). (b) T2-weighted sagital sequence. (c) T2 TIRM sagital sequence. (d) T1/fat sat sagital sequence (T11-L1 segments show hyper intense signal).

Figure 2. (a) T2-weighted axial sequence (segment T11). (b) T2-weighted axial sequence (segment T12). (c) T2- weighted axial sequence (segment T12/L1).
congenital epidermoid cysts originate from displaced ectoderm inclusions arising in early fetal life and may be associated with defective closure of the dural tube [7, 8]. In our case, the patient denied the history of trauma or puncture. Also, it is unreasonable to puncture at such a high level during usual medical procedures. The old age of symptom onset is less possible for a congenital lesion, except for a very slow-growing lesion. Finally, we have supposed that our case was caused either by a congenital lesion with an extraordinary slow-growing pattern or by subtle trauma event with unaware puncture of the cord. Intramedullary epidermoid cysts showed hypointensity on T1-weighted sequence and hyperintensity on T2-weighted sequence. The margins of the lesions had an irregular or a shaggy appearance possibly because of chronic inflammatory response to the squamous tissue leak through the capsule and variable gliosis along the margin [2]. The inner structure of the intramedullary epidermoid cyst was heterogeneous with a high-intensity portion on T2-weighted images. Intravenous injection of gadoteric acid demonstrated peripheral enhancement on T1-weighted images. The acquired cauda equina epidermoid cyst, which is isointense on T1-weighted images and hyperintense on T2-weighted images, reveals faint peripheral enhancement in the post-contrast images [9]. Generally, on MRI examination, the epidermoid cysts have heterogeneous hypointense signal on T1-weighted images and hyperintense signal intensity on T2-weighted images with no contrast enhancement or minimal peripheral enhancement following intravenous gadolinium administration (in our case, the peripheral enhancement was stronger than usual). The differential diagnosis depends on the signal intensity of the content of the cyst and the enhancement pattern of cyst’s wall. However, because of the similar imaging features, it is not easy to make differential diagnosis between epidermoid cyst, dermoid cyst, ventriculus terminalis and arachnoid cyst on MRI examination.

**Figure 3.** (a) T1 fat sat sagital sequence. (b-d) T1 fat sat axial sequence (after administration of intravenous contrast gadoteric acid). The lesion shows minimal peripheral enhancement.

**Figure 4.** Histopathology image analysis shows keratin material characteristic for epidermoid cyst.
tion. Dermoid cysts contain mature tissues of ectoderm with predominance of fatty components, characterized by heterogeneous hyperintensity on all sequences [10]. The high signal intensity on T1-weighted images makes the diagnosis easier as a result of the fatty content of the tumor [11]. The ventriculus terminalis, also known as the fifth ventricle [12], is a small ependyma-lined cavity in the conus medullaris, and it is usually in continuity with the central canal of the rostral spinal cord [13]. The ventriculus terminalis is ovoid, smooth-walled and has no internal septum. The intracystic fluid follows the signals of cerebrospinal fluid (CSF), characterized by low signal on T1-weighted and high signal on T2-weighted sequences. In the eight (72.7%) of 11 cases the contrast gadoteric acid was administered intravenously.

Epidermoid cysts are generally characterized on MRI images by an important variability of signal intensity among different cases and, sometimes, between the different parts of the same cyst; other features include the absence of edema in surrounding tissue, well-defined limits, and the presence of calcifications [14].

Conclusions

Epidermoid cysts should be included in the differential diagnosis when encountering an intramedullary cystic mass with the abovementioned MRI features.

Complete microsurgical excision is the treatment of choice and complete excision or near total excision of this benign tumor can cure the patient with the expectation of nor-
mal life.

**Conflict of Interest**

The authors have no conflict of interest to declare.

**Reference**