Imaging appearance of subependymoma: A rare tumor of the cord

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Abstract

Subependymoma of the cord are rare tumors with very few cases described in the literature. They represent a diagnostic dilemma as far as imaging and histopathology is concerned. They are biologically benign with low proliferative index, hence postoperative prognosis is very good. We present a case of a 42-year-old male patient with an intramedullary Subependymoma located within the central canal of thoracic region cord. On imaging with T1-weighted and T2-weighted sequences it mimicked syringohydromyelia, however, on heavily T2-weighted images the tumor located within the central canal could be delineated. The tumor was excised with complete recovery.

Key words: Cord, magnetic resonance imaging, subependymoma

Introduction

Subependymomas are slow growing tumors usually found in the ventricular system.[1] They possibly account for less than 2% of all spinal cord tumors.[2] On imaging studies they cannot be differentiated from ependymomas[3,4] and can present a diagnostic dilemma. We present a 42-year-old male patient in whom the imaging findings on T1-weighted and T2-weighted Magnetic Resonance (MR) studies mimicked Syringohydromyelia, however on imaging with heavily T2-weighted sequence the tumor located within the central canal could be delineated. Contrast enhanced images did not show any enhancement.

Case Report

A 42-year-old male patient presented with weakness of both lower limbs of two-year duration more on the left side. It was insidious in onset and revealed gradual progression. It progressed to the point when he started having difficulty in getting up from sitting position and developed an ataxic gait. On examination he was found to have spastic paraparesis. Reflexes were increased in both lower. Loss of sensory symptoms was till the inguinal region on left side and up till the knee on right side. On MR Imaging T1 and T2-weighted images revealed dilatation of central canal involving nearly the whole of the thoracic cord, more so from D5-D9 [Figure 1]. The signal intensity of the dilated central canal in the distal cord was slightly different from that of the rest of the central canal suggesting the possibility of a lesion within. No enhancement was seen on contrast enhanced images [Figure 2]. There after T2-weighted 3D TSE sequence was performed, which revealed an ill-defined non-enhancing lesion within the central canal in the region

Figure 1: Sagittal T2W image of the Dorso-Lumbar spine reveals dilatation of the central canal predominantly in the D5-D9 region
of D6-D9 [Figure 3]. It was central in location with left sided predominance. A diagnosis of Subependymoma was given with ependymoma as a differential diagnosis. Patient was thereafter operated upon. Laminectomy was performed from D6-D9. After opening the dura and the arachnoidea, myelotomy was performed extending from the lower part of the thickened cord which was highly vascularized. The tumor was soft reddish gray in color and more on the left side. The tumor was excised along the plane of cleavage and near total removal of the tumor was performed. Following surgery the patient made complete recovery. Histopathology of the excised tumor revealed ependymal cells forming pseudorosettes in a fibrillary background [Figure 4].

Discussion

Spinal subependymomas are much less frequent than their intracranial counterparts, though they become symptomatically obvious quite early. Although Schneiker in 1945 first recognized this variant of ependymoma as a distinct entity, their approximate 40 more cases have been described in the literature and that too in the last two decades. Review of literature performed by Sarkar C et al., revealed a mean age of 47.2 years and male predominance. They also found that the majority of tumors were intramedullary and were located in the cervical region. While Bret et al. found in their review of 29 cases that this tumor accounts for approximately <2% of all spinal cord tumors and that the majority were located in the cervico-thoracic region.

Sarkar et al. on reviewing the literature found no difference on imaging studies between ependymomas and subependymomas. They reviewed the imaging findings of forty cases reported in the literature and found that computed tomography (CT) scan reported in eight cases showed spinal cord enlargement in three and cyst in one. Only two patients showed post contrast enhancement. MRI findings were available for 23 of the 40 reported cases and revealed segmental fusiform dilatation of the cord with low T1-weighted and high T2-weighted signal intensities. Contrast studies were available in 18 of these 23 cases. Ten of 18 cases showed enhancement with contrast, either well circumscribed sharply demarcated areas of homogenous signal enhancement or multiple nodular enhancements. To the best of our knowledge imaging appearance similar to ours has not been described.

These tumors are biologically benign with very low proliferation index. They are eccentrically located within the spinal cord, enabling complete tumor removal in most cases. Surgical removal is usually curative and no further adjunct treatment like radiotherapy should
be given after gross total tumor removal. Radiotherapy given in patients with partial removal of tumor has shown no evidence of any efficacy.\textsuperscript{[12]}

Prognosis is thus determined largely by surgical factors. No recurrence or CSF seedings have been reported till date.\textsuperscript{[4]}

\section*{References}


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