Tzu Chi Medical Journal 26 (2014) 141-143

ELSEVIER

Contents lists available at ScienceDirect

Tzu Chi Medical Journal

journal homepage: www.tzuchimedjnl.com



Case Report

A purely midline ventral schwannoma mimicking a meningioma in the thoracic spine resected via costotransversectomy



Hsiang-Yi Hung^a, Tzu-Yung Chen^{b,c}, Ming-Hsun Li^d, Shin-Yuan Chen^{a,c}, Sheng-Tzung Tsai^{a,c,*}

^a Department of Neurosurgery, Buddhist Tzu Chi General Hospital, Hualien, Taiwan

^b Department of Neurosurgery, Buddhist Tzu Chi General Hospital, Taichung, Taiwan

^c School of Medicine, Tzu Chi University, Hualien, Taiwan

^d Department of Pathology, Buddhist Tzu Chi General Hospital, Hualien, Taiwan

ARTICLE INFO

Article history: Received 1 July 2013 Received in revised form 17 August 2013 Accepted 3 September 2013

Keywords: Schwannoma Spinal cord Surgery

ABSTRACT

Spinal schwannomas are intradural nerve sheath tumors typically located in a dorsolateral or ventrolateral position. Clinical presentations vary from radicular root pain to cord compression myelopathy. Prognosis is usually benign if the tumor can be removed safely. A 62-year-old man had myelopathy and incontinence due to a midline ventral intradural-extramedullary tumor of the thoracic spinal cord. Magnetic resonance imaging demonstrated that the tumor was movable and connected with one root. The tumor was removed with a unilateral costotransversectomy with a posterior approach. Histological diagnosis revealed a schwannoma. Magnetic resonance imaging is a prerequisite to differentiate a ventral and midline intradural spinal schwannoma from other tumors preoperatively. A posterior approach is an efficient and effective method of treating a purely ventral thoracic schwannoma with significant cord compression.

Copyright © 2013, Buddhist Compassion Relief Tzu Chi Foundation. Published by Elsevier Taiwan LLC. All rights reserved.

1. Introduction

Given the typical origin from Schwann cells of the nerve sheath, a spinal schwannoma is one of the most common intraduralextramedullary tumors. A spinal schwannoma almost always locates eccentrically in the dorsolateral, lateral, or ventrolateral position [1]. These tumors can be removed safely through a posterior or posterolateral approach [2]. To our knowledge, a purely ventrally located intradural schwannoma in the thoracic spinal cord is rare [3]. Here, we present a case of a ventral, movable schwannoma of the thoracic spinal cord with myelopathy.

2. Case report

A 62-year-old man presented with progressive numbress and weakness over the bilateral lower limbs for 17 months. He complained of shooting pain with radiation to the bilateral lower limbs

E-mail address: flydream.tsai@gmail.com (S.-T. Tsai).

and also suffered from stool incontinence. Neurological examination revealed bilateral hypesthesia below the T10 dermatome but preserved proprioception. Muscle power in the lower limbs from the bilateral hip flexors to the ankle plantar flexors was graded 3, and the deep tendon reflexes of the knee and ankle were increased. The Babinski sign was positive. Although the voluntary contraction of the anal sphincter was weak, the anal reflex was intact and the anal tone was preserved. Thoracic spine magnetic resonance imaging (MRI) first revealed an isointensity mass located between the T9 and T10 levels over the ventral side of the thoracic cord on T1weighted images (WI; Fig. 1A). In order to differentiate this tumor from other pathological entities, another T1WI scan with contrast enhancement was done 9 days later, and showed a wellencapsulated, homogeneous enhancing mass with a root origin movable over T8 (Fig. 1B). Axial T2WI MRI delineated the margin between the large ventral tumor and the significantly compressed dorsal spinal cord (Fig. 1C).

The patient underwent a right side costotransversectomy at T9 and partial laminectomy at T8. A well-encapsulated tumor was noted after opening the dura (Fig. 1D). The tumor was movable and connected to one nerve root. The tumor was totally removed. After closure of the dura, we inserted transpedicle screws over the right T8, T9, and T10 vertebrae, and applied a rod for fixation.

Conflicts of interest: none.

^{*} Corresponding author. Department of Neurosurgery, Buddhist Tzu Chi General Hospital, 707, Section 3, Chung-Yang Road, Hualien, Taiwan. Tel.: +886 3 8561825x2151; fax: +886 3 8463164.

^{1016-3190/\$ -} see front matter Copyright © 2013, Buddhist Compassion Relief Tzu Chi Foundation. Published by Elsevier Taiwan LLC. All rights reserved. http://dx.doi.org/10.1016/j.tcmj.2013.09.009



Fig. 1. (A) A schwannoma is seen as an isointense mass between the T9–T10 levels over the ventral side of the thoracic cord on T1WI. (B) Another T1 scan with contrast enhancement 9 days later shows a well-encapsulated, homogeneous enhanced mass with a root origin (white arrow) movable over T8. (C) The spinal cord is compressed considerably by the ventral midline schwannoma in axial T2WI. (D) Gross picture of the thoracic spinal schwannoma with root connection (asterisk: tumor; black arrow: root). T1WI = T1-weighted image; T2WI = T2-weighted image.

Somatosensory-evoked potentials were monitored intraoperatively and remained stable. A pathological examination of the tumor revealed a typical schwannoma, which included Antoni A and Antoni B areas and Verocay bodies (Fig. 2). The patient recovered well without complications. His incontinence improved significantly. Two weeks after the operation, muscle power in the lower limbs recovered to grade 4 and he could walk independently.

3. Discussion

Spinal nerve sheath tumors including schwannoma and neurofibroma constitute about one-fourth of intradural spinal tumors and are also the most common intradural extramedullary ones [4]. The majority of spinal schwannomas originate from dorsal sensory roots. Our report with purely midline and ventral located schwannoma might indicate an alternative origin of tumors of Schwann cell derivation [5]. Preoperatively, a schwannoma must be differentiated from a meningioma, another common spinal tumor, which has a more varied location but is often situated anterior to the spinal cord [6]. The initial impression in this case was meningioma. However, most schwannomas are isointense to the spinal cord on T1-weighted images and hyperintense to the cord on T2weighted images [7]. The most important finding was that the second preoperative MRI showed that the tumor moved to a different thoracic level with a root connection, which highlights that MRI can provide significant information for surgical planning.

The location of the spinal tumor may be the most important factor in deciding on the surgical approach. A transthoracic



Fig. 2. Histological samples show typical features of a schwannoma including Verocay bodies in (A) Antoni A (×400) and (B) Antoni B areas (×100).

thoracotomy allows direct visualization of the ventral dura, but extensive mobilization or manipulation of the lungs may lead to significant pulmonary or vascular complications [8]. A lateral extracavitary approach is usually reserved for spinal tumors when extensive paraspinal exposure is required [9]. A costotransversectomy provides access to most ventral intradural spinal lesions, whereas meticulous cord manipulation and varving degrees of bone resection are necessary to prevent endangering neurological function [3]. However, our case experience revealed that the costotransversectomy without destabilization was enough for the ventral intradural-extramedullary movable schwannoma and provided considerable visualization of the tumor. The artery of Adamkiewicz is the largest anterior segmental medullary artery and supplies the spinal cord from T8 to the conus. It arises from the left posterior intercostal artery in 80% of patients, and enters the spinal canal between T9 and T12 in 75% of patients. Therefore, most surgeons advocate a right side approach to avoid inadvertent injury of this artery [10]. The initial presentations of spinal intradural tumors vary in accordance with the level of the tumor. The most common symptom is segmental pain because of direct or indirect irritation of the nerve root or root compression by the tumor [11– 13]. Motor weakness in the lower extremities or incontinence may not be obvious until later stages. However, when tumors locate merely over the ventral side of the spinal cord and compress the spinal cord directly, such as our case, the spinal tracts can be damaged and myelopathy develops as the first clinical presentation [14]. Although a T2 root section could potentially result in Horner syndrome, and lower thoracic motor root division (T8-L1) can cause a painful pseudohernia of the abdominal wall, there is seldom any significant clinical deficit when thoracic nerve roots below T1 are sacrificed [15]. Because the extent of removal is the most significant factor in recurrence, we resected the tumor and the associated root completely [16,17].

In conclusion, although most spinal schwannomas present over the easily accessed dorsolateral position, they should also be considered in a purely midline and ventral thoracic tumor with myelopathy as an initial symptom. Even if the imaging characteristics of a schwannoma usually overlap with meningioma, imaging features such as movability and connected roots are helpful in surgical planning. Unilateral costotransversectomy provides an effective approach for an intradural-extramedullary schwannoma that is ventral and midline to the thoracic spinal cord.

References

- El-Mahdy W, Kane PJ, Powell MP, Crockard HA. Spinal intradural tumours: part I—extramedullary. Br I Neurosurg 1999;13:550–7.
- McCormick PC, Post KD, Stein BM. Intradural extramedullary tumors in adults. Neurosurg Clin N Am 1990;1:591–608.
- [3] Angevine PD, Kellner C, Haque RM, McCormick PC. Surgical management of ventral intradural spinal lesions. J Neurosurg Spine 2011;15:28–37.
 [4] Levy WJ, Latchaw J, Hahn JF, Sawhny B, Bay J, Dohn DF. Spinal neurofibromas:
- [4] Levy WJ, Latchaw J, Hahn JF, Sawhny B, Bay J, Dohn DF. Spinal neurofibromas: a report of 66 cases and a comparison with meningiomas. Neurosurgery 1986;18:331–4.
- [5] O'Toole JE, McCormick PC. Midline ventral intradural schwannoma of the cervical spinal cord resected via anterior corpectomy with reconstruction: technical case report and review of the literature. Neurosurgery 2003;52: 1482–5. discussion 1485–1486.
- [6] Kim MS, Eun JP, Park JS. A dumbbell-shaped meningioma mimicking a schwannoma in the thoracic spine. J Korean Neurosurg Soc 2011;50:264–7.
- [7] Beall DP, Googe DJ, Emery RL, Thompson DB, Campbell SE, Ly JQ, et al. Extramedullary intradural spinal tumors: a pictorial review. Curr Probl Diagn Radiol 2007;36:185–98.
- [8] Lubelski D, Abdullah KG, Steinmetz MP, Masters F, Benzel EC, Mroz TE, et al. Lateral extracavitary, costotransversectomy, and transthoracic thoracotomy approaches to the thoracic spine: review of techniques and complications. J Spinal Disord Tech 2011;26:222–32.
- [9] McCormick PC. Surgical management of dumbbell and paraspinal tumors of the thoracic and lumbar spine. Neurosurgery 1996;38:67–74.
- [10] Steck JC, Dietze DD, Fessler RG. Posterolateral approach to intradural extramedullary thoracic tumors. J Neurosurg 1994;81:202–5.
- [11] Conti P, Pansini G, Mouchaty H, Capuano C, Conti R. Spinal neurinomas: retrospective analysis and long-term outcome of 179 consecutively operated cases and review of the literature. Surg Neurol 2004;61:34–43. discussion 44.
- [12] Celli P, Trillo G, Ferrante L. Spinal extradural schwannoma. J Neurosurg Spine 2005;2:447–56.
- [13] Subaciute J. Early diagnosis of spinal cord schwannoma: the significance of the pain syndrome. Medicina (Kaunas) 2002;38:1086–8.
- [14] Nambiar M, Kavar B. Clinical presentation and outcome of patients with intradural spinal cord tumours. J Clin Neurosci 2012;19:262–6.
- [15] Kim P, Ebersold MJ, Onofrio BM, Quast LM. Surgery of spinal nerve schwannoma. Risk of neurological deficit after resection of involved root. J Neurosurg 1989;71:810–4.
- [16] Seppala MT, Haltia MJ, Sankila RJ, Jaaskelainen JE, Heiskanen O. Long-term outcome after removal of spinal schwannoma: a clinicopathological study of 187 cases. J Neurosurg 1995;83:621–6.
- [17] Ryu KS, Lee KY, Lee HJ, Park CK. Thoracic intramedullary schwannoma accompanying by extramedullary beads-like daughter schwanommas. J Korean Neurosurg Soc 2011;49:302–4.