

Case Report

Solitary Fibrous Tumor One Year After Posterior Cervical Laminectomy



Amir Mahabadi¹, Majid Rezvani¹, Mehdi Shafiei¹, Mehdi Mahmoodkhani¹, Navid Askariardehjani¹

1. Department of Neurosurgery, School of Medicine, Kashani Hospital, Isfahan University of Medical Sciences, Isfahan, Iran



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ABSTRACT

Background and Importance: Cervical laminectomy and fixation is a surgery used to treat symptomatic patients who are resistant to medical therapy. However, it can have serious complications affecting patient outcomes.

Case Presentation: This case report describes a rare occurrence of a solitary fibrous tumor at the site of a previous cervical laminectomy in a 45-year-old female patient. Following non-surgical treatment for paresthesia and weakness in the upper limbs due to cervical stenosis, the patient underwent cervical laminectomy from C3-C6 without instrumentation.

Conclusion: One year after the surgery, the patient presented with severe pain and swelling at the surgical site, and radiological evaluation revealed a large mass in the para-vertebral muscular layer. The mass was removed and the microscopic evaluation revealed a typical solitary fibrous tumor. The patient underwent an extension of laminectomy and fusion with lateral mass screw, and follow-up at 6 months showed no recurrence of the tumor.

* Corresponding Author:

Amir Mahabadi, MD

Address: Department of Neurosurgery, School of Medicine, Kashani Hospital, Isfahan University of Medical Sciences, Isfahan, Iran

Tel: +98 (913) 3623128

E-mail: amirmahabadi@med.mui.ac.ir



Highlights

- Posterior cervical laminectomy can lead to rare soft tissue tumor formation.
- A 45-year-old female developed a solitary fibrous tumor 1 year after surgery.
- The radiological evaluation helped determine the diagnosis of a solitary fibrous tumor.
- The tumor was removed and the patient underwent an extension of laminectomy and fusion.
- Recurrence of the tumor was not observed after a 6-month follow-up.
- Injury to mesenchymal tissue may be a risk factor for solitary fibrous tumors.

Plain Language Summary

This article describes a rare case of a tumor that grew after a patient underwent surgery for a spinal cord problem. The patient was a 45-year-old woman who had surgery to relieve numbness and weakness in her upper limbs. The surgery went well and she was discharged after 3 days. However, a year after the surgery, she returned to the hospital with severe pain and swelling at the surgical site. Further examination revealed a large tumor at the site. The tumor was removed. But the analysis showed it was a tumor called solitary fibrous tumor, a rare type of cancer that can occur in many parts of the body.

1. Background and Importance

Cervical laminectomy and fixation (PCF) is one type of surgery used to treat symptomatic patients resistant to medical therapy; however, this method has serious complications that can affect patient outcomes [1]. Some of the serious complications that can occur after a posterior cervical approach include cerebrospinal fluid (CSF) leakage, surgical site infection (especially when surgery is accompanied by instrumentation), axial pain, fusion failure, graft subsidence, and epidural hematoma [2]. Although the development of a soft tissue tumor after operation is uncommon, this disorder has been reported in the literature [3-6]. Table 1 lists the number of case studies published in this field.

2. Case Presentation

A 45-year-old woman referred to Ayatollah Kashani Hospital, Isfahan, with a major complaint of paresthesia and weakness in the upper limb 2 years ago. Cervical magnetic resonance imaging (MRI) showed spondylosis and moderate to severe cervical stenosis at C3-C6 segments (see pre-operative cervical MRI in Figure 1). Non-surgical treatment methods, such as medication, physical therapy, and cervical collars began following

diagnosis. After 3 months, a follow-up examination showed no recovery. Accordingly, she was a candidate for the operation. The evaluation considered that the appropriate approach was cervical laminectomy from C3-4-5-6 without instrumentation. After general anesthesia, intubation with a GlueScope and prone position, safe laminectomy with a high-speed drill, and decompression of the spinal cord (posterior approach) were performed. The patient was discharged 3 days after the operation in good condition. Post-surgical examination and follow-up imaging demonstrated improvement in motor and sensory findings. After one year, she was referred for severe pain and swelling at the surgical site. She had no new neurological deficit. The radiological evaluation determined a large mass at the para-vertebral muscular layer. The MRI showed a large mass that was hyper-intense at T1 sequences and iso-intense at T2 sequences rather than the CSF signal (Figures 2, 3, and 4). There was no relationship between this lesion and the neural element or thecal sac. The main diagnosis that we were thinking about included organized hematoma, abscess formation, or pseudo-meningocele. Reoperation was programmed. After prone position and incision of the skin, we observed, contrary to our expectations, a large mass at the muscular layer with a thick outer layer (Figure 5). This lesion had no stickiness to the lamina or neural elements and was removed and



Table 1. Some of the case studies published in the field of soft tissue tumor in different levels/locations of the spinal column

Row	Age (y)	Site	Tumor Pathology	Operation	Sex	Period Between First Operation and Dx of Tumor	Ref
1	54	T2-T7	Desmoid tumor	T4 corpectomy+fusion	F	10 months	[3]
2	58	C5-C7	Desmoid fibromatosis	C5-7 Laminectomy+fusion	M	2 years	[4]
3	87	L3-L4	Gossypiboma	Lumbar laminectomy+fusion	M	19 years	[5]
4	76	L3-L5	Desmoid tumor	L3-5 laminectomy+fusion	M	2 years	[6]



sent to the pathology department and reviewed for detection the accurate diagnosis. Microscopic evaluation of the tumor reported fibroblastic mesenchymal cells with spindle shape cells and staghorn vascular which clearly emphasized the diagnosis of solitary fibrous tumor (the typical type). Because of further operation, we decided to extend laminectomy and fusion with a lateral mass screw. Six months later, the follow-up described no tumor recurrence.

3. Discussion

The solitary fibrous tumor was explained by Wagner for the first time at the end of the nineteenth century and further described in 1931 as a connective tissue neoplasm, such as pleura [7]. Although we might assume it is a rare soft tissue neoplasm, perhaps the developments in diagnostic modalities will change our viewpoint in the future (incidence of 0.2/100 000 annually). The solitary fibrous tumor usually accompanies mesenchymal involvement; therefore, some types of this tumor were previously called hemangiopericytoma [8]. One of the unique patterns of this tumor in histology findings was called a “patternless pattern” because of the distraction of cellular architecture [9]. Because

solitary fibrous tumors originate from mesenchymal tissues, they may occur in any region, such as the pleura, abdomen, head, and neck [5]. Similar to many other tumors based on the proliferation rate (mitosis number, hypercellularity), the solitary fibrous tumor will be classified as a “typical” or “malignant” subtype [10]. One of the most important findings in the immunohistochemical evaluation of this tumor is the staining of the transducer and activator of transcription 6 (STAT6), especially in the malignant type [5]. The diagnosis is considered based on the radiologic evaluation. A computed tomography scan showed soft tissue mass with enhancement in about half of the patients. In MRI, the solitary fibrous tumor appears as a variable signal based on the location and subtypes; however, many of the solitary fibrous tumors are iso/hypo in T1 weighted images and iso/hyper in T2 [9]. The main treatment of solitary fibrous tumor is resection but for malignant types, adjuvant therapy may be considered after the operation. Usually, the average follow-up period is about 4-5 years [11]. The recurrence rate is different in the literature from about 4/110 cases (Dimico study) to 18/56 cases (Hohenforset study) [8]. Although solitary fibrous tumor is rare soft tissue cancer, secondary solitary fibrous tumor after cervical laminectomy have been reported rarely.



Figure 1. Cervical magnetic resonance imaging 3 months after cervical laminectomy





Figure 2. Sagittal T2-weighted magnetic resonance imaging of the mass

Notes: C5/C6 congenital block vertebra was described one year after the primary operation.



Figure 3. Sagittal T1-weighted magnetic resonance imaging of the mass

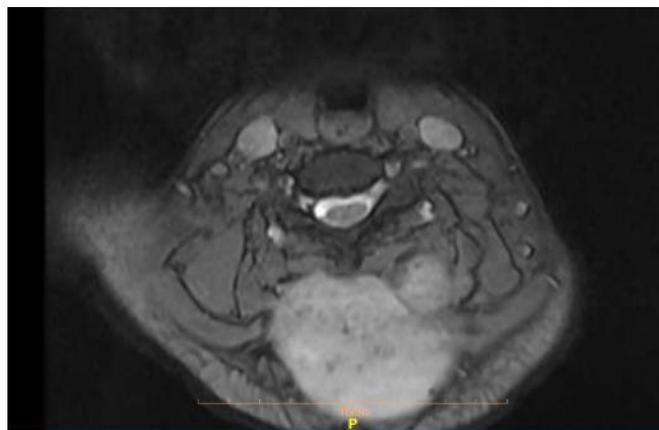


Figure 4. Axial magnetic resonance imaging of the mass



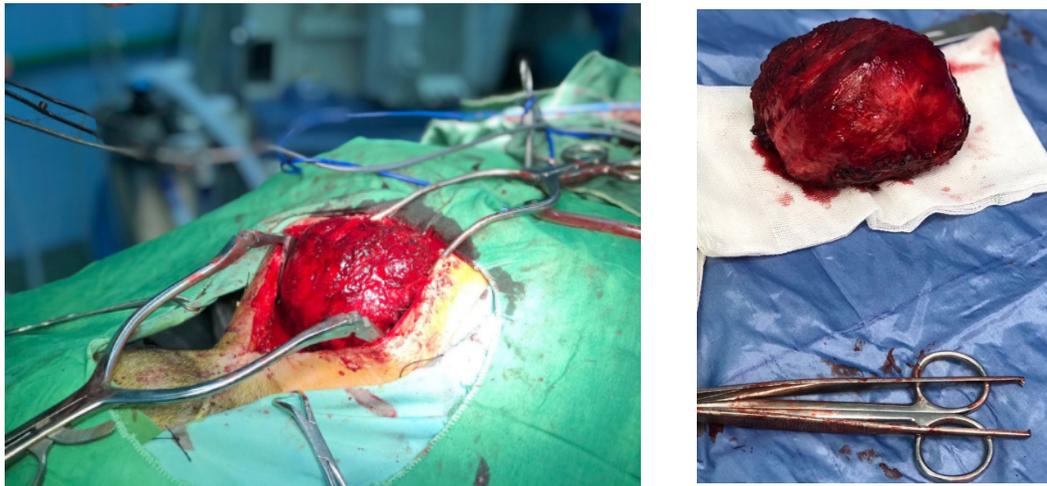


Figure 5. A round mass removed during surgery



4. Conclusion

In this article, we aimed to provide a review of post-operative solitary fibrous tumor development after cervical laminectomy. This case report demonstrated that solitary fibrous tumor at the post-cervical segment is rare. Therefore, after cervical laminectomy, an extremely rare condition underwent tumor resection. Finally, this report states that the mesenchymal tissue injuries can be one of the risk factors for solitary fibrous tumors.

Ethical Considerations

Compliance with ethical guidelines

Written informed consent was obtained from the patient before the operation and for publishing this case report.

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Authors' contributions

Conceptualization and study design: Amir Mahabadi, Majid Rezvani, Mehdi Shafiei and Mehdi Mahmoodkhani; Data collection: Navid Askariardejani; Data analysis and interpretation: Amir Mahabadi; Drafting the article, critically revising and final approval: All authors.

Conflict of interest

The authors declared no conflict of interest.

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References

- [1] Badiee RK, Mayer R, Pennicooke B, Chou D, Mummaneni PV, Tan LA. Complications following posterior cervical decompression and fusion: a review of incidence, risk factors, and prevention strategies. *Journal of Spine Surgery*. 2019; 6(1):323-33. [DOI:10.21037/jss.2019.11.01] [PMID] [PMCID]
- [2] Wang T, Tian XM, Liu SK, Wang H, Zhang YZ, Ding WY. Prevalence of complications after surgery in treatment for cervical compressive myelopathy: A meta-analysis for last decade. *Medicine*. 2017; 96(12):e6421. [DOI:10.1097/MD.0000000000006421] [PMID] [PMCID]
- [3] Puvanesarajah V, Lina IA, Liauw JA, Hsu W, Burger PC, Witham TF. Desmoid tumor formation following posterior spinal instrumentation placement. *Evidence-Based Spine-Care Journal*. 2013; 4(2): 137-42. [DOI:10.1055/s-0033-1357356] [PMID] [PMCID]
- [4] Schlag H, Neuhoff J, Castein J, Hoffmann C, Kandziora F. Sporadic desmoid fibromatosis of the neck after dorsal spondylosis of the cervical spine. *Surgical Neurology International*. 2022; 13:64. [DOI:10.25259/SNI_1240_2021] [PMID] [PMCID]
- [5] Kobayashi T, Miyakoshi N, Abe E, Abe T, Suzuki T, Takahashi M, et al. Gossypiboma 19 years after laminectomy mimicking a malignant spinal tumour: A case report. *Journal of Medical Case Reports*. 2014; 8(1):311. [DOI:10.1186/1752-1947-8-311] [PMID] [PMCID]
- [6] Yan J, Chazen JL. Postoperative lumbar fusion paraspinous desmoid tumor case report. *Radiology Case Reports*. 2022; 17(9):2960-2. [DOI:10.1016/j.radcr.2022.05.022] [PMID] [PMCID]



- [7] Chick JF, Chauhan NR, Madan R. Solitary fibrous tumors of the thorax: Nomenclature, epidemiology, radiologic and pathologic findings, differential diagnoses, and management. *American Journal of Roentgenology*. 2013; 200(3):W238-48. [DOI:10.2214/AJR.11.8430] [PMID]
- [8] Demicco EG, Park MS, Araujo DM, Fox PS, Bassett RL, Pollock RE, et al. Solitary fibrous tumor: A clinicopathological study of 110 cases and proposed risk assessment model. *Modern Pathology*. 2012; 25(9):1298-306. [DOI:10.1038/modpathol.2012.83] [PMID]
- [9] Martin-Broto J, Mondaza-Hernandez JL, Moura DS, Hindi N. A comprehensive review on solitary fibrous tumor: New insights for new horizons. *Cancers*. 2021; 13(12):2913. [DOI:10.3390/cancers13122913] [PMID] [PMCID]
- [10] Baldi, G., et al., Solitary fibrous tumor of all sites: outcome of late recurrences in 14 patients. *Clin Sarcoma Res*. 2013; 3: 4. doi: 10.1186. Europe PMC free article][Abstract][CrossRef][Google Scholar]. [DOI:10.1186/2045-3329-3-4] [PMID] [PMCID]
- [11] Hohenforst-Schmidt W, Grapatsas K, Dahm M, Zarogoulidis P, Leivaditis V, Kotoulas C, et al. Solitary fibrous tumor: A center's experience and an overview of the symptomatology, the diagnostic and therapeutic procedures of this rare tumor. *Respiratory Medicine Case Reports*. 2017; 21:99-104. [DOI:10.1016/j.rmcr.2017.04.007]