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ABOUT COVER

Peer Reviewer of World Journal of Clinical Cases, Suman Baral, MD, Assistant Professor, Department of Surgery, Mediplus Hospital and Trauma Center, Pokhara 33700, Nepal. brylsuman.sur@gmail.com

AIMS AND SCOPE

The primary aim of World Journal of Clinical Cases (WJCC, World J Clin Cases) is to provide scholars and readers from various fields of clinical medicine with a platform to publish high-quality clinical research articles and communicate their research findings online.

WJCC mainly publishes articles reporting research results and findings obtained in the field of clinical medicine and covering a wide range of topics, including case control studies, retrospective cohort studies, retrospective studies, clinical trials studies, observational studies, prospective studies, randomized controlled trials, randomized clinical trials, systematic reviews, meta-analysis, and case reports.

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Retrospective Study

ISSN 2307-8960 (online)

ORIGINAL ARTICLE

Safety and efficacy of posterior approach for resection of spinal meningioma: Impact of dural attachment location

Hong Chen, Ya-Ni Fu, Chu-Di Fu

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Abstract

BACKGROUND

Spinal meningiomas (SMs) are common benign tumors that are typically treated with surgical resection. The choice of surgical approach may vary depending on the location of dural attachment of the SM, with a posterior approach being the traditional preference. However, there is limited research available on the impact of dural attachment location on outcomes following posterior approach for SM resection.

AIM

To investigate the outcomes of posterior approach for SM resection, and compare the results among different dural attachment location subgroups.

METHODS

Between January 2013 and February 2023, a total of 34 SM patients were included in the study. Various clinical and radiologic features, functional states before and after surgery, operating time, intraoperative blood loss, tumor recurrence, and perioperative complications were assessed and compared.

RESULTS

The average age of the included 34 patients' (10 males and 24 females) age was 62.09 years. Mean follow-up duration was 22.65 months. The location of SM was the thoracic spine in 32 cases, with only 2 in the cervical spine. On average, intraoperative blood loss was 520.59 mL, and operating time was 176.76 minutes. Thirty three cases had successful outcomes while only 1 experienced an unexpected outcome. The tumor recurrence rate was 2.9%. After surgery, there were 3 cases of cerebral spinal fluid leakage, 1 case of pneumonia, and 1 case of urinary tract infection. Dural attachments were predominantly found dorsal or dorsolateral (13 cases), followed by ventral or ventrolateral (14 cases), and lateral (7



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Chen H et al. Posterior approach resection of spinal meningioma

cases). The outcomes among these subgroups were similar.

CONCLUSION

The posterior approach for SM resection is safe and effective, yielding comparable surgical and neurological outcomes regardless of the dural attachment location.

Key Words: Spinal meningioma; Posterior approach; Dural attachment; Outcomes; Complications; Recurrence

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Core Tip: This retrospective study aimed to assess the safety and efficacy of the posterior approach for spinal meningiomas resection, and compare the outcomes among different dural attachment location subgroups. Thirty four patients with an average follow-up time of 22.65 months were included. The average operating time was 176.76 min, with intraoperative blood loss of 520.59 mL. Satisfactory outcomes were observed in 97.06% of cases and the tumor recurrence rate was 2.94%. There were no significant differences in operating time, intraoperative blood loss, neurological function, and recurrence rates among three distinct dural attachment location subgroups.

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INTRODUCTION

Spinal meningiomas (SMs) originate from meningothelial arachnoid cap cells and account for 25%-45% of all spinal tumors[1]. These slow-growing intradural extramedullary lesions can develop anywhere along the spine, with a higher prevalence in the thoracic region (67%-84%), followed by the cervical (14%-27%) and lumbar (2%-14%) regions[2]. SMs are most commonly found in women aged between 40 and 70 years, with a female-to-male ratio of 4:1[3].

Fortunately, the majority of SMs are histologically benign or classified as World Health Organization (WHO) grade 1, with only a small percentage considered atypical (WHO grade 2) (5%-25%) or anaplastic (WHO grade 3) (1%-5%)[4]. The most common histological subtypes of SMs include meningothelial, psammomatous, and transitional meningiomas, all of which are classified as WHO grade 1[5]. Patients with SM may present with a range of symptoms from asymptomatic to severe neurological impairments, depending on the extent of spinal cord compression. Typical symptoms include pain, sensory loss, weakness, and bowel or urinary dysfunction[6].

Surgical resection is the primary treatment for SMs, with generally favorable outcomes and a low recurrence rate (1.3%-6.4%)[7]. The choice of surgical approach for resection depends on the tumor site and dural attachment location, typically favoring a posterior approach [8,9]. However, challenges arise when managing with SMs attached ventrally or ventrolaterally. Some surgeons have attempted anterior approaches for these cases but note a higher complication rate[10, 11]. Others argue that posterior approaches can provide sufficient exposure for safe removal of ventral or ventrolateral SMs[12-14]. The optimal method for resecting SMs with ventral or ventrolateral dural attachment remains a topic of ongoing debate.

This study aimed to evaluate the safety and efficacy of using a posterior approach for resecting SMs with different sites and dural attachments locations. Additionally, the study examined whether different dural attachments influenced outcomes.

MATERIALS AND METHODS

Patients

This study retrospectively identified patients who underwent SM resection in the Department of Orthopedics at 903 Hospital of the Joint Logistic Support Force of the People's Liberation Army from January 2013 to February 2023. Inclusion criteria were: SM was confirmed by pathology; the patient underwent a posterior approach; had complete medical records and imaging data. Exclusion criteria were: Presence of other neurogenic tumors; did not undergo a posterior approach; had severe cardiovascular or cerebrovascular disease and could not tolerate anesthesia and surgery. Finally, 34 patients were included in the study.

Data collection

The data collected included information on age, sex, tumor level and length, dural attachment location, preoperative symptoms, duration of symptoms, operative time, intraoperative blood loss, instrumented fusion, perioperative complic-



ations, hospital length of stay, time to follow-up, and recurrence. Additionally, 3-dimensional computed tomography and magnetic resonance imaging (MRI) were used to assess tumor size, calcification, spinal cord compression, and dural attachment location.

The modified McCormick grade (MMG) scale was used to classify neurological function, visual analogue scale (VAS) scores were used to assess pain, the Simpson grading scale was used to grade resection extension, and WHO tumor grade was used for histological evaluation.

Surgical procedure

All patients underwent surgery using the median posterior approach while in the prone position. A partial or total laminectomy was carried out to access the tumor. Subsequently, the tumor was grossly resected after making a midline incision in the dura. The dural attachment was either removed or cauterized using bipolar coagulation forceps. Following this, the dura was either continuously sutured or covered with artificial dura. In cases where total laminectomy was performed, pedicle screw fixation was performed, followed by posterolateral bone graft fusion to stabilize the spine.

Statistical analysis

The data were analyzed using SPSS 25 (IBM Corp., Armonk, NY, United States). Categorical variables were reported as count and percentage, while continuous variables were presented as mean \pm SD. The Kolmogorov-Smirnov test was used to assess normal distribution. Fisher's exact test was used to compare categorical variables, the Student's *t* test to evaluate normally distributed variables between 2 groups, the Mann-Whitney *U* test to compare non-normally distributed variables between 2 groups, and the Kruskal-Wallis H test to compare 3 groups. Statistical significance was set at a *P* value < 0.05.

RESULTS

Patient data

The study included a total of 34 patients, consisting of 24 females and 10 males with a sex ratio of 2.4:1. The average age was 62.09 years (ranging from 27 to 83 years). The average hospital stay was 24.76 days (ranging from 9 to 39 days), and the average follow-up time after surgery was 22.65 months (ranging from 6 to 36 months). Of these cases, 31 patients presented with pain and/or myelopathy, while 3 cases were incidentally found to have meningiomas. Preoperative symptoms included neck/back/radicular pain (23.53%), sensory deficit (73.53%), motor deficit (70.59%), and urinary dysfunction (5.88%), with an average duration of 15.12 months (ranging from 0 to 72 months). SMs were located in the cervical spine in 2 cases and in the thoracic spine in 32 cases. The location of dural attachments of the tumors were 8 dorsal, 5 dorsolateral, 3 ventral, 11 ventrolateral and 7 lateral. Tumor length was less than 1 cm in 2 cases, between 1 and 2 cm in 24 cases, and greater than 2 cm in 8 cases. Patient information is listed in Table 1.

Neurological function and pain were assessed using the MMG and VAS scores respectively. Preoperatively, MMG classifications I, II, III, IV, and V were observed in 4, 2, 6, 14, and 8 cases, respectively, with a mean VAS score of 4.03 (range 0-7) (Table 2).

Surgical resection and clinical outcomes

As illustrated in Table 2, a posterior approach was utilized for all cases, with 11 undergoing partial laminectomy without internal fixation and 23 undergoing total laminectomy with instrumented fusion. Of these cases, 5 underwent Simpson grade 1 tumor resection, while the remaining 29 cases underwent Simpson grade 2 resection. The average operating time was 176.76 minutes (ranging from 80 to 310 minutes), and the average intraoperative blood loss was 520.59 mL (ranging from 100 to 1700 mL). Of these cases, 25 tumors were non-calcified and 9 were calcified. The histological type of 33 cases was WHO grade 1 (meningothelial 17, psammomatous 10, transitional 4, and fibrous 2), with only 1 case classified as WHO grade 2 (atypical).

The distribution of MMG grades postoperatively (1 week after surgery) were as follows: 5 cases were MMG I, 9 cases were MMG II, 12 cases were MMG III, 7 cases were MMG IV, and 1 case was MMG V. At the final follow-up, 26 cases were classified as MMG I, 7 cases as MMG II, and 1 case as MMG III. Satisfactory outcomes, defined as no or minimal function deficit at follow-up (MMG I or II), were observed in 97.06% of cases, while unsatisfactory outcomes, defined as no change in dysfunction or postoperative MMG III-IV, were found in 2.94% of cases. The mean VAS score decreased from 2.03 postoperatively to 0.41 at the final follow-up. All cases showed improvement in neurological function and significant pain relief at the final follow-up compared to preoperative and postoperative assessments (P < 0.05). Perioperative complications included cerebral spinal fluid (CSF) leakage in 3 cases, pneumonia in 1 case, and urinary tract infection in 1 case. Three patients with CSF leakage had drainage tubes placed in the surgical area for 5-7 days to ensure adequate CSF drainage; 1 patient with pneumonia and 1 patient with urinary tract infection received antibiotics to control the infection. All complications were resolved before discharge, and had no long-term effects on patient outcomes. Recurrence occurred 9 years after surgery in a 27-year-old woman with a tumor located ventrolateral to the spinal cord in the cervical spine, presenting with a long dural tail. The initial surgery involved Simpson grade 1 resection followed by radiotherapy. The recurrence rate was 2.94%.

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Table 1 Patient information

Case number	Age (years)	Sex	Clinical presentation	Duration of symptoms (months)	Spinal segment	Dural attachment location	Calcification	Tumor length (cm)
1	68	Male	Pain; Sensory and motor deficit	4	T8	Ventrolateral	No	1.66
2	73	Male	No	0	T8/9	Dorsolateral	Yes	1.88
3	71	Female	Pain	24	T4	Dorsolateral	Yes	2.26
4	60	Female	Pain; Sensory deficit	12	T7-9	Dorsolateral	Yes	1.66
5	57	Female	No	0	T9-11	Ventrolateral	No	2.00
6	68	Female	Pain; Sensory and motor deficit	24	T4	Ventrolateral	Yes	1.35
7	54	Female	Sensory and motor deficit	5	T9/10	Ventral	No	1.69
8	45	Male	No	0	T12	Lateral	No	1.57
9	61	Female	Pain; Sensory deficit	12	T11/12	Dorsal	Yes	1.63
10	69	Female	Sensory and motor deficit	12	T3	Ventral	No	1.70
11	58	Female	Sensory and motor deficit; Urinary dysfunction	72	T6/7	Ventrolateral	Yes	1.66
12	63	Female	Sensory and motor deficit	24	T11/12	Dorsal	No	2.63
13	69	Female	Sensory and motor deficit	4	T5/6	Ventrolateral	No	1.61
14	53	Male	Sensory and motor deficit	12	T9/10	Dorsal	No	1.26
15	71	Female	Sensory and motor deficit	12	C5/6	Lateral	Yes	1.74
16	57	Male	Pain; Sensory deficit	12	T5/6	Lateral	No	1.39
17	58	Female	Sensory and motor deficit	12	T7/8	Lateral	No	1.46
18	76	Female	Sensory and motor deficit	2	T7/8	Dorsolateral	No	1.20
19	71	Male	Sensory and motor deficit	60	T4/5	Dorsal	No	1.34
20	58	Female	Sensory and motor deficit	36	T7/8	Ventrolateral	No	2.22
21	50	Male	Sensory and motor deficit	6	T7/8	Dorsal	No	1.46
22	49	Female	No	0	T10/11	Dorsal	No	1.44
23	72	Female	Sensory and motor deficit	24	Т9	Ventral	No	1.39
24	73	Female	Motor deficit; Urinary dysfunction	12	T7/8	Lateral	No	1.54
25	50	Female	Sensory and motor deficit	12	T4	Ventrolateral	No	0.94
26	56	Female	Pain; Sensory and motor deficit	3	T3/4	Ventrolateral	No	2.51
27	68	Male	Sensory deficit	2	T4	Dorsal	No	2.25
28	70	Female	Pain; Sensory deficit	48	T8/9	Dorsal	Yes	1.93
29	58	Female	Sensory and motor deficit	1	T10/11	Ventrolateral	Yes	1.56
30	61	Male	Sensory and motor deficit	12	T1/2	Ventrolateral	No	1.45
31	27	Female	Sensory and motor deficit	7	C3-6	Ventrolateral	No	4.05
32	83	Male	Motor deficit	12	T6/7	Dorsolateral	No	1.60
33	71	Female	Motor deficit	12	T11/12	Lateral	No	2.50
34	63	Female	Sensory and motor deficit	24	T5/6	Lateral	No	0.80

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Incidental 3 Incidental 15.2 ± 16.86 (0-72) Duration of symptoms (months) 16.86 ± 7.91 (0-3.90) Follow-up duration (months) 26.5 ± 12 (6-3.61) Operative time (minutes) 16.76 ± 5.0.22 (80-31.01) Operative blood loos (mL) 25.0 ± 38.0.41 (100-170.01) Calification 25.0 ± 38.0.41 (100-170.01) Calification 25.0 ± 38.0.41 (100-170.01) No 25.0 ± 38.0.41 (100-170.01) Statiogical types 25.0 ± 38.0.41 (100-170.01) Yes 9 No 25.0 ± 38.0.41 (100-170.01) Statiogical types 25.0 ± 38.0.41 (100-170.01) Histological types 26.0 ± 38.0.41 (100-170.01) Statiogical types 26.0 ± 38.0.41 (100-170.01) Statiogical types 10 Stational 20.0 ± 38.0.41 (100-170.01) Stational 20.0 ± 38.0.41 (100-170.01) Stational 20.0 ± 38.0.41 (100-170.01) Stational 20.0 ± 38.0.41 (100-100.01) Stational 20.0 ± 38.0.41 (100-100.01) Stational 20.0 ± 38.0.41 (100-100.01) Stational <t< td=""><td>Symptoms</td><td></td></t<>	Symptoms			
İvation of symptoms (months) 15.2 ± 16.8 (0.472) Hoppial length of stay (days) 0.8 ± 7.9 (0.493) Follow-up duration (months) 26.5 ± 12.0 (0.433) Operative time (minutes) 16.0 ± 12.0 (0.433) Intaoperative blood loos (m.l.) 20.5 ± 38.0 ± 10.0 ±	Pain/myelopathy	31		
<table-container><table-container><table-container><table-container><table-container><table-container><table-container><table-container><table-container><table-container><table-container><table-container><table-container><table-container><table-container><table-container><table-container><table-row><table-row><table-row><table-row><table-row><table-row><table-row><table-row><table-row><table-row><table-row><table-row><table-row><table-row><table-row><table-row><table-row><table-row><table-row><table-row><table-row></table-row></table-row></table-row></table-row></table-row></table-row></table-row></table-row></table-row></table-row></table-row></table-row></table-row></table-row></table-row></table-row></table-row></table-row></table-row></table-row></table-row></table-container></table-container></table-container></table-container></table-container></table-container></table-container></table-container></table-container></table-container></table-container></table-container></table-container></table-container></table-container></table-container></table-container>	Incidental	3		
Follow-up duration (months)2.65 ± 2 (6-36)Operative information767 ± 5 - 22 (80-310)Intaoperative blood loss (mL)20.5 ± 380.41 (100-1700)Claffication2No2Yes9Stabulary Constraints9Fistological types10Pannonatous0Fistonational2Fistonational2Stabulary Constraints10Stabulary Constraints2Fistonational2Arastional2Augusta2Augusta3Augusta3Augusta3Augusta1Augusta3 <td>Duration of symptoms (months)</td> <td>15.12 ± 16.86 (0-72)</td>	Duration of symptoms (months)	15.12 ± 16.86 (0-72)		
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CacificationNo5Yes9Histological types7Meningothelial10Snamomatous4Fransitional2Atypical2NtO grade1132132Simong rege1	Operative time (minutes)	176.76 ± 50.22 (80-310)		
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Hitological typesMeningothelia70Psannomatous0Transitional4Forous2Atypical0WHO grade11332131Storous or use1Storous or use1 <t< td=""><td>No</td><td>25</td></t<>	No	25		
Meningothelial17Psannomatous0Transitional4Fibrous2Atypical0WHO grade113321Support grade1	Yes	9		
Psanmonatous 10 Transitional 4 Fibrous 2 Atypical 1 VHO grade 33 2 1 Supson grade 1	Histological types			
Transitional4Fibrous2Atypical1WHO grade313321Simpson grade	Meningothelial	17		
Fibrous2Atypical1WHO grade321Sinson grade-	Psammomatous	10		
Atypical 1 WHO grade 3 1 32 2 1 Simpson grade 5	Transitional	4		
WHO grade 1 33 2 Janoba 33 Simpson grade	Fibrous	2		
1 33 2 1 Simpson grade	Atypical	1		
2 1 Simpson grade	WHO grade			
Simpson grade	1	33		
	2	1		
1 5	Simpson grade			
	1	5		

Chen H et al. Posterior approach resection of spinal meningioma

2	29
Instrument fusion	
No	11
Yes	23
Perioperative complications	
CSF leakage	3
Pneumonia	1
Urinary tract infection	1
Recurrence	1

MMG: Modified McCormick grade; WHO: World Health Organization; CSF: Cerebral spinal fluid.

Comparison of dorsal/dorsolateral, ventral/ventrolateral and lateral dural attachment subgroups

Patients with dorsal or dorsolateral dural attachment were categorized as the dorsal/dorsolateral subgroup (n = 13), those with ventral or ventrolateral dural attachment were categorized as the ventral/ventrolateral subgroup (n = 14), and those with lateral dural attachment were categorized as the lateral subgroup (n = 7). There were no significant differences in age, sex, preoperative MMG and VAS scores, degree of tumor calcification, Simpson grades, and follow-up duration between these subgroups (P > 0.05), indicating that the data among the 3 subgroups were comparable. Following surgery and at the final follow-up, improvements in neurological function and pain relief were observed in all subgroups. Additionally, the MMG and VAS scores, operative time, intraoperative blood loss, perioperative complications, and recurrence rates were similar across the 3 subgroups (P > 0.05) (Table 3).

Representative cases

Two representative cases are presented in Figure 1 and Figure 2. Case 1 involved a 68-year-old woman who had been experiencing pain, numbness, and weakness in both lower limbs for 2 years. Prior to surgery, her neurological function was assessed as MMG IV and she had a VAS score of 6. Imaging revealed a calcified tumor in the thoracic spine with ventrolateral dural attachment measuring 1.35 cm in length. The patient underwent a posterior approach for total laminectomy and a Simpson grade 2 resection. The surgery lasted 230 minutes with an intraoperative blood loss of 1200 mL. Pedicle screw fixation and posterolateral bone graft fusion were performed for spinal stabilization. The histological type of the tumor was WHO grade 1 (psammomatous). There were no perioperative complications during her hospital stay. Postoperatively, the patient's MMG score improved to III and VAS score decreased to 3. At the final follow-up 3 years after surgery, the patient's MMG score was I and VAS score was 0 (Figure 1).

Case 2 involved a 58-year-old female patient who presented with pain and numbness in both lower limbs, along with difficulty walking steadily for a month. Prior to surgery, her neurological function was classified as MMG III and she reported a VAS score of 4. MRI results indicated a tumor in the thoracic spine with ventrolateral dural attachment, measuring 1.56 cm in length. The surgical procedure involved a posterior approach for partial laminectomy and a Simpson grade 2 resection, lasting 180 minutes with intraoperative blood loss of 600 mL. Histological analysis revealed a WHO grade 1 (psammomatous) tumor type. There were no complications during the patient's hospital stay. Postoperatively, the patient's MMG score improved to II and VAS score decreased to 1. At the final follow-up 2 years post-surgery, the MMG score was grade I and VAS score was 0 (Figure 2).

DISCUSSION

Among the 34 patients analyzed in this study, 70.59% were female. The majority of patients exhibited symptoms such as local/radicular pain, sensory/motor deficit, and urinary tract dysfunction. Most SMs were found in the thoracic spine and were non-calcified, with prevalent histological types being meningothelial, psammomatous, and transitional. The epidemiology, clinicoradiologic characteristics, and histological types observed in this study closely resembled those documented in previous reports[15].

The primary treatment for symptomatic SMs was maximal surgical resection, with adjuvant treatments including radiotherapy and chemotherapy. All patients in this study underwent Simpson grade 1 or 2 resection, resulting in good functional and neurological outcomes during the follow-up period. Only 1 patient experienced relapse 9 years postsurgery. Sarikaya et al[16] reported recurrence in 2 patients under 18 years old with cervical SMs and long dural tails after Simpson grade 2 resection [16]. These patients underwent Simpson grade 1 resection upon recurrence and remained in remission. The authors suggested that young patients with cervical SMs and long dural tails might be at higher risk of recurrence. The characteristics of the recurrent patient in our study aligned with those reported by Sarikaya et al[16] supporting their hypothesis. Misra et al[17] proposed a classification system for SMs and emphasized the importance of instrumented fusion to prevent delayed spinal deformity or instability post-tumor excision[17]. Total laminectomy with facet joint resection was identified as a predictor of such issues, with patients typically undergoing laminoplasty with



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Table 3 Comparison of dorsal/dorsolateral, ventral/ventrolateral and lateral dural attachment subgroups, mean ± SD (range)						
Variable	Dorsal/dorsolateral (<i>n</i> = 13)	Ventral/ventrolateral (n = 14)	Lateral (n = 7)	P value		
Age (years)	65.23 ± 10.35	58.93 ± 11.37	62.57 ± 10.1	0.314		
Sex						
Male	6	2	2	0.208		
Female	7	12	5			
Preoperative MMG						
I/II/III	6	4	2	0.996		
IV/V	7	10	5			
Postoperative MMG						
I/II/III	12	9	5	0.495		
IV/V	1	5	2			
Final follow-up MMG						
I/II/III	12	11	4	0.208		
IV/V	1	3	3			
Preoperative VAS score	3.54 ± 2.67	4.64 ± 1.6	3.71 ± 2.81	0.732		
Postoperative VAS score	1.62 ± 1.33	2.57 ± 1.09	1.71 ± 1.25	0.11		
Final follow-up VAS score	0.38 ± 0.65	0.5 ± 0.65	0.29 ± 0.49	0.738		
Calcification						
No	8	11	6	0.538		
Yes	5	3	1			
Operative time (min)	185.38 ± 53.29	174.29 ± 33.62	165.71 ± 73.68	0.751		
Intraoperative blood loss (mL)	569.23 ± 449.79	478.57 ± 254.74	514.29 ± 491.35	0.8		
Simpson grade						
1	4	1	0	0.189		
2	9	13	7			
Perioperative complications						
CSF leakage	2	1	0			
Urinary tract infection	0	1	0	0.602		
Pneumonia	0	0	1			
Follow-up duration (months)	19.31 ± 12.53 (6-36)	24 ± 11.6 (6-36)	26.14 ± 12.08 (6-36)	0.34		
Recurrence						
No	13	13	7	1		
Yes	0	1	0			

MMG: Modified McCormick grade; VAS: Visual analogue scale.

micro-titanium plate fixation for spinal stability. Posterolateral fusion with pedicle screw fixation was also utilized in our study, with no instances of internal fixation failure or spinal deformity during follow-up. Following the dural opening and subsequent repair, CSF leakage was the most common complication, with a reported incidence ranging from 0% to 4%. Additional perioperative complications can include pneumonia, urinary tract infection, surgical site infection, spinal cord edema, deep venous thrombosis, pulmonary embolism and myocardial infarction. CSF leakage, pneumonia and urinary tract infection were observed in our study, with the incidence of CSF leakage consistent with previous literature [18]. Spinal cord edema is a prevalent complication that can occur following spinal cord decompression. Clinically, this condition is characterized by deterioration in neurological function rather than improvement. To mitigate or reduce the discomfort associated with spinal cord edema, we administered glucocorticoids during surgery after spinal cord decompression, followed by a regimen of glucocorticoids and nerve dehydration medications for 2-3 days postoper-

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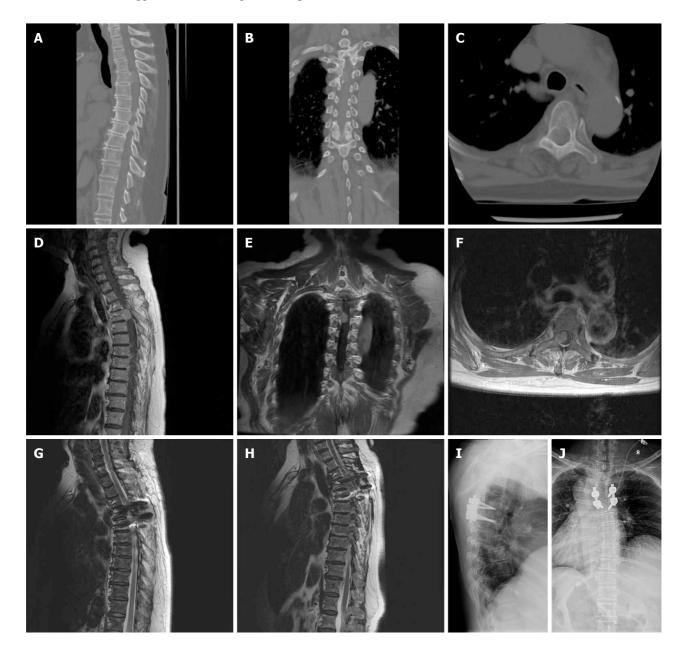


Figure 1 Imaging finding in case 1. A: Sagittal; B: Coronal; C: Axial 3-dimensional CT images revealed calcification within the tumor; D: Subsequent sagittal; E: Coronal; F: Axial magnetic resonance imaging T1-weighted post-contrast-enhanced images demonstrated homogeneous contrast enhancement of the tumor, which was located in the thoracic spine with ventrolateral dural attachment; G: Sagittal magnetic resonance imaging (MRI) T2-weighted image displayed gross-total resection of the tumor; H: Sagittal MRI T2-weighted image displayed no recurrence of the tumor observed 3 years after surgery; I: Lateral; J: Anteroposterior X-ray images depicted pedicle screw fixation and posterolateral bone graft fusion procedures that were conducted to stabilize the spine.

atively.

SMs are often detected due to symptoms of spinal cord compression, but some cases are found incidentally. MRI is considered the gold standard for detection, with SMs appearing isointense to the spinal cord on T1-weighted images and isointense or hypointense on T2-weighted images. T1-weighted post-contrast-enhanced images typically show homogeneous enhancement in SMs[19]. Characteristic radiological findings in SMs include the 'dural tail' and 'gingko leaf' signs on contrast-enhanced MRI[20]. Management options for asymptomatic SMs usually involve close observation or surgical resection with patient consent. In this study, three incidental cases opted for surgical removal of their tumors after providing informed consent.

The most direct approaches for SMs resection were based on the dural attachment locations, which include anterior, lateral, and posterior approaches. SMs in the cervical spine are best approached anteriorly *via* corpectomy for removal, but dural repair can be challenging in this scenario. For SMs with anterior or lateral dural attachments at T3-L2, a lateral extracavitary approach or costotransversectomy can be utilized for excision. Posterior fixation may be necessary with these approaches due to extensive pedicle removal and facetectomy, with attention given to the great vessels and radicular arteries. While the posterior approach has traditionally been the preferred method for SM resection, the safety and efficacy of this approach for SMs with ventral or ventrolateral dural attachments are still under debate. In our study, some surgeons believed that they were familiar with the posterior approach and this approach could successfully remove

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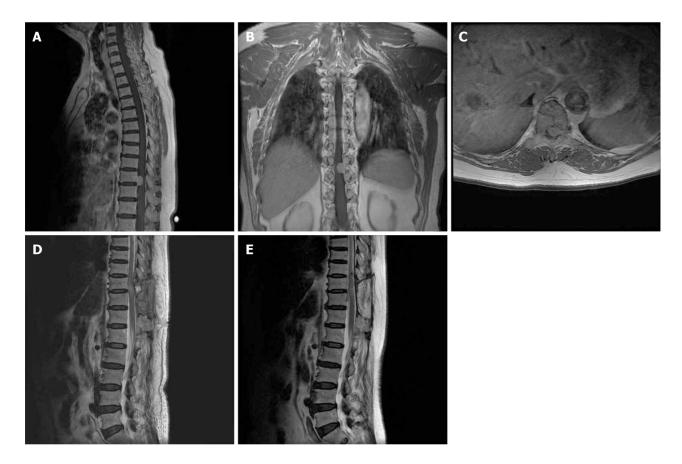


Figure 2 Imaging finding in Case 2. A: Sagittal; B: Coronal; C: Axial magnetic resonance imaging (MRI) T1-weighted post-contrast-enhanced images revealed a homogeneously contrast-enhanced tumor located in the thoracic spine with ventrolateral dural attachment; D: Sagittal MRI T2-weighted image indicated gross-total resection of the tumor; E: Sagittal MRI T2-weighted image indicated no recurrence of the tumor observed 2 years after surgery.

the tumor; thus, some patients with SMs on the ventral or ventrolateral side were selected to undergo the posterior approach. The results from this study show that there were no significant differences in outcomes among patients with dorsal/dorsolateral, ventral/ventrolateral and lateral dural attachments who underwent the posterior approach for resection, indicating that the posterior approach may be adequate for any dural attachments.

Although a myriad of new technological advances such as surgical microscopes, intraoperative ultrasound, ultrasonic tumor aspirators, intraoperative neurological monitoring and microscope-based augmented reality have been used to enhance the resection of SMs and minimize neural tissue injury [21,22], many hospitals do not have these new medical devices. Our retrospective study suggested that posterior approach resection of SMs accompanied by detailed preoperative planning, meticulous operation and strict postoperative management might be a common, safe and effective method for SM treatment.

Given the retrospective nature of this study, it is important to note the limitations such as the lack of extensive followup for certain patients due to changes in medical providers and electronic medical records. Moreover, all patients in the study were selected to undergo the posterior approach by their respective surgeons, which may have introduced a selection bias. Additionally, the small sample size restricts the ability to draw robust conclusions on the safety and efficacy of resecting SM with any dural attachment location through a posterior approach.

Despite these constraints, this study stands out as one of the few that investigated the outcomes of patients with SM and compares these outcomes across various dural attachment subgroups in individuals who underwent the posterior approach for resection. It is recommended that future studies should include prospective trials, with multi-center collaborations and larger patient cohorts with longer follow-up durations.

CONCLUSION

SMs are benign tumors with favorable prognoses after surgical resection. However, resecting SMs with a ventral or ventrolateral dural attachment using a posterior approach can be challenging. The impact of dural attachment location on outcomes following posterior resection of SMs remains poorly understood. This study found that posterior resection of SMs with various dural attachment locations resulted in good outcomes, with no significant differences in neurological outcomes, Simpson grade, complications, or recurrence rates among the different subgroups. These findings highlight the feasibility of successfully resecting any SM through via posterior approach, with consistently positive outcomes regardless of the dural attachment location.



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FOOTNOTES

Author contributions: Chen H provided the idea, designed the study and wrote the manuscript; Fu YN collected and analyzed the data, and prepared the tables and figures; Fu CD revised the manuscript and supervised the study. All authors have read and approved the final manuscript.

Institutional review board statement: The study was reviewed and approved by No. 903 Hospital of PLA Joint Logistic Support Force.

Informed consent statement: The studies involving humans were approved by the Ethics Committee of 903 Hospital and conducted in accordance with local legislation and institutional requirements. Signed consent was waived because the study was a retrospective review of medical records and involved the preservation of anonymity during data collection, statistical analysis and manuscript writing.

Conflict-of-interest statement: The authors declare that the study was conducted in the absence of any commercial or financial relationships, and there is no conflict of interest.

Data sharing statement: All data can be obtained from the first author.

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