Original Article

Correlation of Intraoperative Neurophysiological Parameters and Outcomes in Patients with Intramedullary Tumors

Abstract

Introduction: Intramedullary spinal cord tumors (IMSCT) account for about 2%-4% of all central nervous system tumors. Surgical resection is the main treatment step, but might cause damage to functional tissues. Intraoperative neuromonitoring (IONM) is an adopted measure to decrease surgical complications. Below, we describe the results of IMSCT submitted to surgery under IONM at a tertiary institution. **Methods:** The sample consisted of consecutive patients with IMSCT admitted to the Neurological Institute of Curitiba from January 2007 to November 2016. A total of 47 patients were surgically treated. Twenty-three were male (48.9%) and 24 were female (51.1%). The mean age was 42.77 years. The mean follow-up time was 42.7 months. Results: Neurological status improved in 29 patients (62%), stable in 6 (13%), and worse in 12 (25%). Patients who presented with motor symptoms at initial diagnosis had a worse outcome compared to patients with sensory impairment and pain (P = 0.026). Patients with a change in electromyography had worse neurological outcomes compared to patients who did not show changes in monitoring (P = 0.017). Discussion and Conclusion: No prospective randomized high evidence study has been performed to date to compare clinical evolution after surgery with or without monitoring. In our sample, surgical resection was well succeeded mainly in oligosymptomatic patients with low preoperative McCormick classification and no worsening of IONM during surgery. We believe that microsurgical resection of IMSCT with simultaneous IONM is the gold standard treatment and achieved with good results.

Keywords: Intraoperative neurophysiology, outcome, spinal cord tumors, surgery

Introduction

Intramedullary spinal cord tumors (IMSCT) comprise the least common types of spinal neoplasms. They account for about 2%-4% of all central nervous system tumors and 20%-30% of all spinal cord tumors.[1-10] The most common type of IMSCT is ependymoma, astrocytoma.[11-22] bv followed manifested as back pain, radicular pain, or neuropathic pain has been reported to be the most common presenting symptom in patients with an IMSCT. Other common symptoms include motor disturbances and sensory symptoms (dermatomal, saddle, or segmental). Sphincter disturbances may occur as an early symptom.[11-22]

Without treatment, IMSCT can lead to severe neurologic deterioration with serious motor deficits, including paraplegia or even quadriplegia. The most important part of the treatment is surgical resection. Radical resection has been associated with increased long-term overall survival. [23-28]

This is an open access journal, and articles are distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 4.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms.

For reprints contact: WKHLRPMedknow reprints@wolterskluwer.com

However, surgery might cause damage to functional tissues, which leads to neurologic complications.^[25,26]

To reduce iatrogenic the risk of complications, intraoperative neuromonitoring (IONM) is a worldwide adopted measure.[18-28] Several different monitoring modalities are currently in use.[8-10] Although IONM is widely used, and many single-center studies have been conducted, no high evidence study has clearly defined the added value, in terms of overall sensitivity and specificity to detect or prevent neuronal injury of the different monitoring techniques used in IMSCT surgery.[16-28]

Below, we describe the results of IMSCT submitted to surgery under IONM at a tertiary institution. Clinical and radiological data, lesion features, timing of symptom onset, and IONM findings were recorded. The IONM included continuous needle electromyography.

How to cite this article: Ruschel LG, Aragão A, Oliveira MF, Milano JB, Neto MC, Ramina R. Correlation of intraoperative neurophysiological parameters and outcomes in patients with intramedullary tumors. Asian J Neurosurg 2021;16:243-8.

Submitted: 14-May-2020 **Revised:** 09-Aug-2020 **Accepted:** 05-Oct-2020 **Published:** 28-May-2021

Leonardo Gilmone Ruschel^{1,2,3}, Afonso Aragão¹, Matheus Fernandes de Oliveira^{2,3,4}, Jerônimo Buzetti Milano¹, Mauricio Coelho Neto¹, Ricardo Ramina¹

¹Department of Neurosurgery, Neurological Institute of Curitiba, Curitiba, Paraná, ²DFV Neuro, ³Hospital Alemão Oswaldo Cruz, ⁴Department of Neurosurgery, Hospital do Servidor Público Estadual de São Paulo, São Paulo, Brazil

Address for correspondence:
Dr. Matheus Fernandes de
Oliveira,
Rua Loefgren 700, Apto 103,
Vila Mariana, São Paulo,
Brazil. ZIP CODE 04040-000.
E-mail: mafernoliv@yahoo.
com.br



We evaluated the outcome according to the modified McCormick scale.

Methods

The study sample consisted of consecutive patients with IMSCT admitted to the Neurological Institute of Curitiba (INC) Neurosurgical Ward from January 2007 to November 2016. Patients enrolled in the study were then followed during treatment. We evaluated age, gender, symptoms, histology, surgical data (IONM), and postoperative outcome.

Inclusion criteria

All patients undergoing microsurgery as a treatment strategy for IMSCT were eligible for inclusion.

Criteria for surgery

Criteria used to indicate surgery were as follows: patients with intramedullary spinal cord lesions, incidental or symptomatic. Patients in palliative care and/or with very low functional status (Karnofsky score <40) were not considered candidates for surgery.

Signs and symptoms were described according to neurological impairments: pain, sensorial disturbances, reflexes, motor strength, gait and balance disturbances, and sphincterian changes. Patients were classified according to McCormick classification (ranging from I to IV). Postoperatively, patients were then compared and considered with neurological improvement, stability, or neurological impairment.

Sample

A total of 47 patients were surgically treated. Twenty-three were male (48.9%) and 24 were female (51.1%). The mean age was 42.77 years, ranging from 7 to 80 years. The mean follow-up time was 42.7 months (ranging from 8 to 120 months). Survival analysis was extracted from the INC registry of patient data.

Surgical approach

In all cases, the institutional routine is to perform laminotomy of target levels and midline myelotomy with microsurgical technique. The surgical aim is to perform maximal safe resection without neurological impairment.

Neuromonitoring evaluation

IOM was recorded continuously, from before patient positioning, at which point baseline signals were obtained, until waking up from anesthesia. Our evaluation was limited to continuous electromyography. We did not apply somatosensory evoked potential (SSEPs), motor evoked potentials (MEPs), D-waves, or dorsal column mapping. Signals were obtained from all four extremities.

Statistical analysis

Statistical analyses were performed using SPSS Statistics, IBM, Armonk, New York, USA. We applied the Chi-square

test when applicable. We considered statistically significant data when P < 0.05.

Ethics

The study protocol was approved by the Institutional Ethics in Research Board.

Results

Location of tumors

Cervical tumors occurred in 23 patients (40.4%), while cervicothoracic in 3 (6.4%) patients and thoracic in 25 (53.2%) patients. The extension of tumor was correspondent to 2 levels or less in 37 (78%) subjects, while 3 or more in 10 subjects (22%).

Gross total resection was achieved in 40 patients (85%), while partial resection was obtained in 7 patients (15%).

Histology

As described in Table 1, the most common tumor was ependymoma, with 23 cases (48.9%), followed by hemangioblastomas (6 cases, 12.8%) and astrocytomas (5 cases, 10.6%). We had also atypical pathological specimens, such as chordoma and lymphoma. There were two cases of metastases, being both of melanomas. We illustrate pre- and postoperative images of typical cases in Figure 1.

Table 1: Histological types			
	n (%)		
Cavernoma	4 (8.5)		
Astrocytoma	5 (10.6)		
Arachnoid cyst	2 (4.3)		
Chordoma	1 (2.1)		
Ependymoma	23 (48.9)		
Ganglioglioma	1 (2.1)		
Hemangioblastoma	6 (12.8)		
Lymphoma	1 (2.1)		
Arteriovenous malformation	1 (2.1)		
Neurocysticercosis	1 (2.1)		
Melanoma metastasis	2 (4.3)		
Total	47 (100)		

Table 2: Initial symptoms in evaluated patients			
Symptom	n (%)		
Sensorial	29 (62)		
Pain	28 (60)		
Motor	15 (32)		
Hyperreflexia	5 (11)		
Sphincter	5 (11)		
Gait	6 (13)		
Asymptomatic	2 (4)		



Figure 1: Pre- and postoperative images of typical cases. Above, pre- and postoperative magnetic resonance of an ependymoma. Below, pre- and postoperative magnetic resonance of a hemangioblastoma

Clinical presentation

The initial clinical presentation is presented in Table 2. The main symptoms were sensorial (29 cases, 62%), followed by pain (28%–60%) and motor deficits (15 cases, 32%). Hyperreflexia, esfincterian, and gait symptoms completed the main symptoms. Two patients were symptom free at the diagnosis.

Preoperative McCormick

Preoperative McCormick was I in 8 subjects (17%), II in 22 subjects (46.8%), III in 15 patients (32%), and IV in 2 other patients (4.2%).

Follow-up

The mean follow-up time was 42.7 months (ranging from 8 to 120 months). In the late follow-up (1 year), neurological status was improved in 29 patients (62%), stable in 6 (13%) patients, and worse in 12 (25%). Overall survival rate free of disease was 90%, with three cases of postoperative complications (6%), being one case of pulmonary thromboembolic event, and two cases of cerebrospinal fluid fistula. At the end of the follow-up, we diagnosed one case of melanoma central nervous system dissemination and one case of tumoral progression instead of surgery.

Intraoperative neuromonitoring parameters and correlation to outcome

We divided IONM parameters into three groups: a group without decrease in intraoperative electromyography, a second group with decrease and recovery during surgery, and the last group with decrease without recovery.

In the late follow-up (1 year), neurological status was improved in 29 patients (62%), stable in 6 (13%) patients, and worse in 12 (25%). Among 47 patients, 24 (51%) did not present intraoperative signal decrease, while 23 (49%) presented decrease, being 12 patients (25.5%) with recovery and 11 (23.5%) without recovery.

Among 29 patients with clinical improvement, 19 had no decrease in IONM (65%), while 10 had decrease (35%), being 9 with recovery and 1 without. Among 6 patients clinically stable, 3 had no decrease in IONM (50%), while 3 had decrease (50%), being 0 with recovery and 3 without. Among 12 patients clinically worse, 2 had no decrease in IONM (8%), while 10 had decrease (92%), being 3 with recovery and 7 without [Table 3].

Table 4 illustrates parameters evaluated and postoperative functional outcome. Gender, mean age, histology, anatomical localization, extension, preoperative McCormick, and resection rate did not change the outcome. Factors involved in outcome included initial motor symptoms presentation and intraoperative change in IONM. The worse outcome occurred in male patients, above 40-year-old, with partial resection and McCormick III or IV.

Patients who presented with motor symptoms at initial diagnosis had a worse outcome compared to patients with sensory impairment and pain (P=0.026). Patients with a change in electromyography (reduction in peak amplitude >50%, increase in peak latency >10%, and total loss of waveform) had worse neurological outcomes compared to patients who did not show changes in monitoring (P=0.017). Patients who showed changes in monitoring and returned to baseline potential during surgery had better neurological outcomes compared to patients who did not return to baseline wave pattern (P=0.001) [Table 4].

Discussion

Intramedullary tumors account for 2%–4% of central nervous system tumors and 10% of spinal tumors, with ependymomas and astrocytomas being the most frequent histological types. [1-10] These tumors are usually benign, slow growing and may extend to various segments of the spinal cord and have few specific symptoms because they usually evolve insidiously. [11-16] Even with the progress in surgical technique and intraoperative monitoring, postoperative prognostic factors are conflicting in the literature.

Ependymomas are common in adults, while astrocytomas are far more common in children.^[10] Most ependymomas have relatively demarcated borders, while astrocytomas are more infiltrative and need to be resected until a white matter "interphase" appears.^[20]

Microsurgical resection of IMSCTs is currently considered the primary treatment modality, while radiotherapy and/or chemotherapy are reserved for recurrent or malignant tumors.^[3,9] The observation that the majority of IMSCTs are benign and consequently gross-total removal might result in long-term survival further supports the need for safe resection.^[10] The extent of resection has been correlated with progression-free survival and lower recurrence

Table 3: Correlation of intraoperative neuromonitoring parameters and clinical outcome

Tubic 5: Colli	ciation of intraoperat	ive near omomitoring parame	ters and enimear outco	me
	Total, <i>n</i> (%)	Improvement, n (%)	Stable, n (%)	Worse, n (%)
IONM	47	29 (61.7)	6 (12.7)	12 (25.5)
No decrease	24 (51.1)	19 (65)	3 (50)	2 (17)
Decrease	23 (48.9)	10 (35)	3 (50)	10 (83)
Decrease with recovery	12 (25.5)	9 (31)	0	3 (25)
Decrease without recovery	11 (23.4)	1 (4)	3 (50)	7 (63)

IONM-Intraoperative neuromonitoring

Table 4: Factors related to postoperative outcome									
Factor	Total , <i>n</i> (%)	Improvement, n (%)	Stable, n (%)	Worsening, n (%)	P				
Gender									
Male	23 (48.9)	13 (56.5)	2 (8.7)	8 (34.8)	0.318				
Female	24 (51.1)	16 (66.7)	4 (16.7)	4 (16.7)					
Age									
<40	20 (42.6)	13 (65.0)	3 (15.0)	4 (20)	0.735				
>40	27 (57.4)	16 (59.3)	3 (11.1)	8 (29.6)					
Histology									
Neuroepithelial	29 (61.7)	18 (62.1)	4 (13.8)	7 (24.1)	0.941				
Not neuroepithelial	18 (38.3)	11 (61.1)	2 (11.1)	5 (27.8)					
Location									
Cervical	19 (40.4)	12 (63.2)	2 (10.5)	5 (26.3)	0.747				
Cervicothoracic	3 (6.4)	2 (66.7)	1 (33.3)	0 (0)					
Thoracic	25 (53.2)	15 (60)	3 (12.0)	7 (28.0)					
Extension (levels)									
<3	37 (78.7)	20 (54.1)	5 (13.5)	12 (32.4)	0.083				
≥3	10 (21.3)	9 (90)	1 (10)	0 (0)					
IONM change									
Yes	23 (48.9)	10 (43.5)	3 (13.0)	10 (43.5)	0.017				
No	24 (51.1)	19 (79.2)	3 (12.5)	2 (8.3)					
IONM									
Improvement	12 (52.2)	9 (75.0)	0 (0)	3 (25.0)	0.001				
No improvement	11 (47.8)	1 (9.1)	3 (27.3)	7 (63.6)					
McCormick									
I and II	30 (63.8)	19 (63.3)	5 (16.7)	6 (20)	0.365				
III and IV	17 (36.2)	10 (58.8)	1 (5.9)	6 (35.3)					
Extent of resection									
Total	40 (85.1)	27 (67.5)	4 (10)	9 (22.5)	0.133				
Partial	7 (14.9)	2 (28.6)	2 (28.6)	3 (42.9)					
Symptoms									
Pain	28 (59.6)	20 (71.4)	4 (14.3)	4 (14.3)	0.026				
Motor	15 (31.9)	9 (60)	2 (13.3)	4 (26.7)					
Sensorial	29 (61.7)	19 (65.5)	4 (13.8)	6 (20.7)					

IONM-Intraoperative neuromonitoring

rates.^[21] Advances in microsurgery have contributed to safer resection ability. The most common surgical complication is postoperative cerebrospinal fluid (CSF) leak which occurs in about 5%–10% of surgeries.^[12,14]

However, despite all advances, surgery for IMSCTs is still very challenging and may carry significant morbidity. The most significant risk associated with surgery for IMSCTs is spinal cord injury and a resulting neurological deficit. Neurophysiological monitoring is an important tool intended to eliminate or reduce this

surgical risk.^[16-28] To date, both SSEP and transcranial MEP monitoring techniques have been used. Continuous electromyography and D-wave are also recommended, increasing the sensitivity and specificity of intraoperative findings.^[16-28]

No prospective randomized high evidence study has been performed to date to compare clinical evolution after surgery with or without monitoring. [6,28] A recent meta-analysis failed to find a strong support for IONM use in spinal surgery. [23] On the other hand, there was a slight

tendency to support IONM in IMSCT surgery.^[23] Thus, the use of IONM is still only supported series of low scientific evidence. In addition, there are no studies comparing different modalities of IONM in IMSCT and thus, there is no established superiority of one modality over the other.^[16-28]

Even though, the routine neurosurgical practice has incorporated IONM and the advocated advantages include detecting positioning disturbances, intraoperative disturbances due to anesthetic and surgical intervention, and it adds information to a neurosurgeon when deciding to continue resection or stopping it, avoiding further tissue damage. [6-14] Factors associated with improved outcomes for IMSCT include preoperative neurological status, presence of a tumor dissection plane, tumor size, the use of neuromonitoring, and postoperative radiation therapy. [6,11,12]

In our sample of 47 patients, histology was similar to most series in the literature, prevailing ependymomas, astrocytomas, and hemangioblastomas. Symptoms were mainly sensorial, pain, and motor symptoms. Surgical resection was the best treatment mainly in oligosymptomatic patients with low preoperative McCormick classification (I and II) and no worsening of IONM during surgery. Worst results were observed in patients with preoperative motor symptoms, high McCormick classification (III and IV), and with decrease in IONM sign and persistent decrease in the postoperative period.

There are some limitations of the study. At first, the small sample size does not allow us to perform an individualized analysis concerning each histological type. Then, we did not apply routinely SSEPs and MEPs and D-wave monitoring. We applied exclusively continuous electromyography which might decrease the final power to identify monitoring changes and thus interfere with optimal results. However, once there is not still a consensus on the ideal monitoring setup, we believe that this fact may interfere but not invalidate our results.

We believe that microsurgical resection of IMSCT is the gold standard treatment and achieved with acceptable complications. Simultaneous use of IONM is for us mandatory, mitigating positioning and manipulation problems and taken into account when deciding surgical and resection extent goals. The ideal setup of which monitoring modalities in each case is yet to be delimitated.

Financial support and sponsorship

Nil.

Conflicts of interest

There are no conflicts of interest.

References

1. Hyun SJ, Rhim SC. Combined motor and somatosensory evoked potential monitoring for intramedullary spinal cord tumor

- surgery: Correlation of clinical and neurophysiological data in 17 consecutive procedures. Br J Neurosurg 2009;23:393-400.
- Quiñones-Hinojosa A, Lyon R, Zada G, Lamborn KR, Gupta N, Parsa AT, et al. Changes in transcranial motor evoked potentials during intramedullary spinal cord tumor resection correlate with postoperative motor function. Neurosurgery 2005;56:982-93.
- Korn A, Halevi D, Lidar Z, Biron T, Ekstein P, Constantini S. Intraoperative neurophysiological monitoring during resection of intradural extramedullary spinal cord tumors: Experience with 100 cases. Acta Neurochir (Wien) 2015;157:819-30.
- Ghadirpour R, Nasi D, Iaccarino C, Giraldi D, Sabadini R, Motti L, et al. Intraoperative neurophysiological monitoring for intradural extramedullary tumors: Why not? Clin Neurol Neurosurg 2015;130:140-9.
- Rijs K, Klimek M, Scheltens-de Boer M, Biesheuvel K, Harhangi BS. In reply to the letter to the editor regarding "Intraoperative neuromonitoring in patients with intramedullary spinal cord tumor: A systematic review, meta-analysis, and case series". World Neurosurg 2019;127:664.
- Cannizzaro D, Mancarella C, Nasi D, Tropeano MP, Anania CD, Cataletti G, et al. Intramedullary spinal cord tumors: The value of intraoperative neurophysiological monitoring in a series of 57 cases from two Italian centres. J Neurosurg Sci 2019 Sep 23. doi: 10.23736/S0390-5616.19.04758-1.
- Nagoshi N, Tsuji O, Nakashima D, Takeuchi A, Kameyama K, Okada E, et al. Clinical outcomes and prognostic factors for cavernous hemangiomas of the spinal cord: A retrospective cohort study J Neurosurg Spine 2019;31:271-8.
- Persson O, Fletcher-Sandersjöö A, Burström G, Edström E, Elmi-Terander A. Surgical treatment of intra- and juxtamedullary spinal cord tumors: A population based observational cohort study. Front Neurol 2019;10:814.
- Sun XY, Wang W, Zhang TT, Kong C, Sun SY, Guo MC, et al. Factors associated with postoperative outcomes in patients with intramedullary Grade II ependymomas: A Systematic review and meta-analysis. Medicine (Baltimore) 2019;98:e16185.
- Khalid S, Kelly R, Carlton A, Wu R, Peta A, Melville P, et al. Adult intradural intramedullary astrocytomas: A multicenter analysis. J Spine Surg 2019;5:19-30.
- 11. Benjamin CG, Frempong-Boadu A, Hoch M, Bruno M, Shepherd T, Pacione D. Combined use of diffusion tractography and advanced intraoperative imaging for resection of cervical intramedullary spinal cord neoplasms: A case series and technical note. Oper Neurosurg (Hagerstown) 2019;17:525-30.
- 12. Hamilton KR, Lee SS, Urquhart JC, Jonker BP. A systematic review of outcome in intramedullary ependymoma and astrocytoma. J Clin Neurosci 2019;63:168-75.
- Rijs K, Klimek M, Scheltens-De Boer M, Biesheuvel K, Harhangi BS. Intraoperative neuromonitoring in patients with intramedullary spinal cord tumor: A systematic review, meta-analysis, and case series. World Neurosurg 2019;125:498-510.e2.
- Goyal A, Yolcu Y, Kerezoudis P, Alvi MA, Krauss WE, Bydon M. Intramedullary spinal cord metastases: An institutional review of survival and outcomes. J Neurooncol 2019;142:347-54.
- Scibilia A, Terranova C, Rizzo V, Raffa G, Morelli A, Esposito F, et al. Intraoperative neurophysiological mapping and monitoring in spinal tumor surgery: Sirens or indispensable tools? Neurosurg Focus 2016;41:E18.
- Verla T, Fridley JS, Khan AB, Mayer RR, Omeis I. Neuromonitoring for intramedullary spinal cord tumor surgery. World Neurosurg 2016;95:108-16.
- 17. Cheng JS, Ivan ME, Stapleton CJ, Quinones-Hinojosa A,

- Gupta N, Auguste KI. Intraoperative changes in transcranial motor evoked potentials and somatosensory evoked potentials predicting outcome in children with intramedullary spinal cord tumors. J Neurosurg Pediatr 2014;13:591-9.
- 18. André-Obadia N, Mauguière F. Electrophysiological testing in spinal cord tumors. Neurochirurgie 2017;63:356-65.
- Siller S, Szelényi A, Herlitz L, Tonn JC, Zausinger S. Spinal cord hemangioblastomas: Significance of intraoperative neurophysiological monitoring for resection and long-term outcome. J Neurosurg Spine 2017;26:483-93.
- Ghadirpour R, Nasi D, Iaccarino C, Romano A, Motti L, Sabadini R, et al. Intraoperative neurophysiological monitoring for intradural extramedullary spinal tumors: Predictive value and relevance of D-wave amplitude on surgical outcome during a 10-year experience. J Neurosurg Spine 2018;30:259-67.
- Barzilai O, Lidar Z, Constantini S, Salame K, Bitan-Talmor Y, Korn A. Continuous mapping of the corticospinal tracts in intramedullary spinal cord tumor surgery using an electrified ultrasonic aspirator. J Neurosurg Spine 2017;27:161-8.
- Lakomkin N, Mistry AM, Zuckerman SL, Ladner T, Kothari P, Lee NJ, et al. Utility of Intraoperative Monitoring in the Resection of Spinal Cord Tumors: An Analysis by Tumor Location and Anatomical Region. Spine (Phila Pa 1976) 2018;43:287-94.

- Daniel JW, Botelho RV, Milano JB, Dantas FR, Onishi FJ, Neto ER, et al. Intraoperative Neurophysiological Monitoring in Spine Surgery: A Systematic Review and Meta-Analysis. Spine (Phila Pa 1976) 2018;43:1154-60.
- Kumar N, G V, Ravikumar N, Ding Y, Yin ML, Patel RS, et al. Intraoperative Neuromonitoring (IONM): Is There a Role in Metastatic Spine Tumor Surgery? Spine (Phila Pa 1976) 2019;44:E219-24.
- Taricco MA, Guirado VM, Fontes RB, Plese JP. Surgical treatment of primary intramedullary spinal cord tumors in adult patients. Arq Neuropsiquiatr 2008;66:59-63.
- Guirado VM, Taricco MA, Nobre MR, Couto EB Jr, Ribas ES, Meluzzi A, et al. Quality of life in adult intradural primary spinal tumors: 36-Item Short Form Health Survey correlation with McCormick and Aminoff-Logue scales. J Neurosurg Spine 2013;19:721-35.
- Novak K, Widhalm G, de Camargo AB, Perin N, Jallo G, Knosp E, et al. The value of intraoperative motor evoked potential monitoring during surgical intervention for thoracic idiopathic spinal cord herniation. J Neurosurg Spine 2012;16:114-26.
- Deletis V, Bueno De Camargo A. Interventional neurophysiological mapping during spinal cord procedures. Stereotact Funct Neurosurg 2001;77:25-8.