Original Article

Ann Rehabil Med 2017;41(3):413-420 pISSN: 2234-0645 • eISSN: 2234-0653 https://doi.org/10.5535/arm.2017.41.3.413



Contrast Spread in the Superoposterior Approach of Transforaminal Epidural Steroid Injections for Lumbosacral Radiculopathy

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Objective To observe the contrast spread in superoposterior transforaminal epidural steroid injection (SP TFESI) and investigate the correlation between spread patterns and efficacy.

Methods Thirty-one patients with lumbosacral radiculopathy underwent single-level TFESI under fluoroscopy. The final needle tip position was targeted toward the SP quadrant of the intervertebral foramen. To observe the spread, 1 mL of contrast material was injected, followed by a steroid injection. The contrast spread was graded anteroposteriorly and vertically in the epidural space. The effect of SP TFESI was evaluated by proportional pain score reduction.

Results Levels injected were L4-5 (n=20) and L5-S1 (n=11). Seventeen cases were lateral, and 14 were central herniated disc (HD). Baseline mean visual analog scale score was 6.23. Contrast dispersed dorsally in all the cases, and 45.2% cases showed a concurrent ventral spread. The proportion of the pain reduction after 2 weeks showed no difference between the two groups. In vertical spreading analysis, mean cephalic/caudal grades were 1.40/1.55 at L4-5 level and 1.73/1.64 at L5-S1 level. The HD location had no effect on contrast dispersion.

Conclusion In SP TFESI, ventral contrast spread did not guarantee a better effect; however, the extent of cephalic flow in ventral expansion group correlated with the proportion of pain reduction.

Keywords Epidural injection, Contrast media, Lumbosacral, Radiculopathy, Treatment outcome

INTRODUCTION

Epidural steroid injection (ESI) is a widely used interventional treatment for low back pain caused by lumbosacral radiculopathy [1,2]. Among the ESI techniques, the trans-

Received June 13, 2016; Accepted October 3, 2016

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foraminal approach is considered as a preferred method for effective delivery of the injectate to the ventral epidural space, close to the dorsal root ganglion [1-4]. Conventionally, when transforaminal epidural steroid injections (TFESI) are administered, the needle is placed below

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the underside of the pedicle and above the exiting spinal nerve root, within the upper half of the neural foramen, namely called as 'safe triangle.' Specifically, the target of the needle is the superoanterior (SA) quadrant of the neural foramen. However, this conventional technique occasionally encounters the following problems. (1) Because the needle must traverse the nerve root to reach the target, chances of touching the nerve root is highly probable. (2) In a severely degenerated and narrowed foramen, accessing the epidural space is very hard [5]. Furthermore, since the radicular artery (the artery of Adamkiewicz) are most commonly in the SA quadrant of the foramen [6-9], these arteries can be punctured, causing catastrophic complications [10,11]. Therefore, an alternative technique has been proposed in which the target of the needle is superoposterior (SP) quadrant of the foramen [5,12-14]. A previous study reported that the ventral flow and extent of contrast medium spreading resulting in SA TFESI [15]. However, in SP TFESI, because the needle is placed at the posterior aspect of the foramen, it is unclear whether the medication can reach the ventral epidural space and relieve pain [5].

This study aimed to observe the contrast spread pattern in SP TFESI and investigate the correlation between the spread pattern and clinical efficacy. In addition, the relationship between the anatomical pathology and contrast spread patterns in SP TFESI was analyzed.

MATERIALS AND METHODS

Subjects

This study was conducted from November 2014 to December 2015. We recruited 31 subjects diagnosed with lumbar radiculopathy caused by a herniated intervertebral disc, undergoing TFESI. Each patient complained of persistent low back pain and radicular pain in the lower extremity for at least 2 weeks despite appropriate medication and conservative treatment. Diagnosis was confirmed by physical examination, electrophysiological study, and magnetic resonance (MR) images. Patients <18 years, >80 years, or pregnant were excluded from the study. Other exclusion criteria were as follows: (1) history of previous lumbar surgery, (2) presence of systemic inflammatory disease, (3) uncontrolled diabetes, (4) allergy to contrast medium or lidocaine, and (5) injections within 3 months. Patients were allowed to take medications prescribed before the study; however, additional pain medications were not prescribed during the study.

This study was approved by the regional Institutional Review Board, and written informed consents were obtained from all the patients.

MR image analysis

The primary pathology causing radiculopathy was determined according to the report of the radiologists of the author's hospital and classified as central herniated disc (HD) or lateral HD for further analysis of the structures. In accordance with the 'zone system' [16], central HD was defined as the HD medial to the medial edge of the articular facets, and lateral HD was defined as the HD lateral to the medial edge of the articular facets.

Procedure

All epidural injections were administered in an outpatient procedure room by one of two physicians with extensive experience in fluoroscopically guided lumbar TFESI. The criteria for choosing the candidate target levels were determined by the HD location of the MR images and clinical symptoms as follows: (1) Patients with lateral HD at L4-5 level on MR images underwent L4-5 TFESI. (2-1) Patients with central HD at L4-5 level and symptoms of L5 radiculopathy underwent L5-S1 TFESI. (2-2) In case of difficulty in approaching the L5-S1 foramen despite central HD at L4-5 level, TFESI was performed using the L4-5 foramen instead. Some studies [17,18] and our own clinical experience reported a successful therapeutic effect with TFESI performed by substituting the foramen one level above. (3) The patients with lateral and central HD at L5-S1 level and symptom of L5 radiculopathy underwent L5-S1 TFESI.

Subjects were placed in the prone position on a fluoroscopy table, and the skin was prepared in a sterile manner with an iodine antiseptic solution and draped. The Carm was set to optimally visualize the target foramen, rotated in the range 20° - 25° to the lesion side to provide an oblique view, and tilted cranially or caudally at 0° - 5° if needed to align the vertebral end plates. The overlying soft tissue was anesthetized using 2% lidocaine HCl (Huons, Hwaseong, Korea). A 23-gauge, 3.5-inch spinal needle with a curved tip was advanced into the region of the involved nerve root under fluoroscopic guidance. As this study aimed to evaluate only SP TFESI, the final

needle tip position was targeted towards the SP quadrant of the intervertebral foramen, where the needle was placed under the median inferior edge of the pedicle on oblique view, and anterior to the posterior margin of the intervertebral foramen on lateral view. To confirm the exact injection position and observe the pattern of spread, 1 mL of contrast material Xenetix 300 (Guerbet, Gorinchem, The Netherlands) was injected. Furman et al. [19] demonstrated that when 1 mL of contrast (ISOVUE 320; Bracco Diagnostics, Princeton, NJ, USA) was used on SA TFESI, the contrast reached the superior aspect of the superior intervertebral disc (IVD) or inferior aspect of the inferior IVD in 46% of the total attempts, for each landmark. Thus, we assumed that 1 mL of contrast would also be sufficient to observe the contrast spread in SP TFESI. After the needle was confirmed in the SP quadrant position without intravascular flow or soft tissue infiltration and the contrast spread was assessed, steroidal drug was injected. The drug composed of 2 mL of 0.5% lidocaine, 0.5% lidocaine HCl (Samjin, Hwaseong, Korea), and 20 mg (0.5 mL) of triamcinolone acetonide (Triam; Shinpoong, Seoul, Korea) and was slowly administered.

Measurements

General characteristics such as age, sex, height, weight, and body mass index of all the patients were evaluated. Prior to the TFESI, the baseline pain score was evaluated using the visual analog scale (VAS). At 2 weeks after TFE-SI, the follow-up VAS score was measured. To compare the clinical efficacy according to the spreading pattern, the percentage of the pain relief was calculated using the following formula: (baseline pain score – follow-up pain score)×100/baseline pain score.

Ventral and dorsal contrast spread patterns were analyzed according to the following criteria. Ventral spreading, contrast medium reached the posterior aspect of the vertebral body or intervertebral disc; dorsal spreading, contrast medium reached the lamina or ligamentum flavum. In addition, vertical spread patterns for the cephalic and caudal directions were separately evaluated. A grading system was used for the extent of contrast spread to a predetermined position by adopting Furman's concept [19], emphasizing the importance of various relevant pathologic landmarks in lumbar radiculopathy. The IVD is important to a central HD. As the opening of the dural sleeve spans the upper half of the pedicles at L3, L4, and L5 vertebrae, the vertical midpoint of the pedicle was also considered to be an important landmark for the entrance between the central canal and spinal root (Fig. 1). The grading system was as follows (Fig. 2): Grade 0, no visible spread; Grade 1, within the same vertebral level; Grade



Gr 4 Pedicle Gr 3 Nerve root Superior IVD Gr 2 Gr 1 0 pt _____Injection level Gr 1 Inferio Gr 2 IVD Gr 3 Gr 4 **E**pidural space Caudal

Vertebra

Fig. 1. CT myelography showing L3, L4, and L5 nerve roots. Openings of the dural sleeves span the upper half of the pedicles at L3, L4, and L5 vertebrae. CT, computed tomography.



Cephalic

2, within the vertical level of the superior IVD (cephalad) or inferior IVD (caudad); Grade 3, above the superior IVD, not exceeding the horizontal midline of the upper level pedicle, or below the inferior IVD, not exceeding the horizontal midline of the lower level pedicle; Grade 4, over the horizontal midline of the upper or lower pedicle. The grade of spreads was separately evaluated in cephalic and caudal directions (Fig. 3). In the case of dorsal isolated spread, the grade was derived from the dorsal spread (ventral expansion), the grade was derived from the highest one between the ventral and dorsal spreads.

Statistical analysis

The distribution of anteroposterior patterns of contrast spread was examined according to the isolated ventral, dorsal, or concurrent spread. Using the odds ratio (OR), the differential likelihood of anteroposterior contrast spread at each injection level was determined. Student t-test was used to compare the treatment effect in each group showing different contrast spread according to the ventral and dorsal sides of the epidural space. Pearson correlation analysis was performed to reveal the association between the treatment effect and extent of vertical contrast spread. Fisher exact test was used to investigate the correlation between the lesion location (lateral or central HD) and contrast spread. Statistical analyses were performed using the SPSS software package ver. 12.0 (SPSS Inc., Chicago, IL, USA). The p-values <0.05 were considered to indicate statistical significance.

RESULTS

General characteristics of all the patients are listed in Table 1. TFESI was performed at the L4-5 and L5-S1 levels in 20 and 11 patients, respectively. The main pathology at the injected level was lateral HD in 17 patients and central HD in 14. Fifteen patients had a lateral HD at the L4-5 level on MR images and underwent L4-5 TFESI. Of these, only one patient was confirmed to have L4 radiculopathy by an electrophysiologic study, and five had L5 radiculopathy. The others had normal electrophysiologic findings, but complained of symptoms of L5 radiculopathy. Five patients with central HD at L4-5 level were suspected to have L5 radiculopathy, but underwent L4-5 TFESI because of anatomical changes in the lumbosacral area: 3 cases with a high iliac crest, 1 case of a large L5 transverse process, and 1 case of pseudo-arthrosis of L5 vertebra with the iliac crest. Six patients with central HD at L4-5 level and 2 with lateral HD at L5-S1 level underwent L5-S1 TFESI. Three patients had central HD at L5-S1 level, but presented symptoms suggesting L5 radiculopathy and underwent L5-S1 TFESI (Supplement Table S1 shows clinical presentations, the results of MRI and EMG,



Fig. 3. Anteroposterior (A) and lateral (B) fluoroscopic view of L5-S1 SP TFESI. Contrast filled the dorsal epidural space and spread to the ventral space. Vertical dispersal analysis showed Grade 3 (above the superior IVD, not exceeding the horizontal midline of the upper level pedicle) in the cephalic direction and Grade 2 (within the vertical level of the inferior IVD) in the caudal direction. SP TFESI, superoposterior transforaminal epidural steroid injection; IVD, intervertebral disc.

and ESI-performed level for each patient). The mean VAS score of all the patients was 6.23±1.33 at baseline.

The contrast medium spread dorsally in all cases (n=31), and 45.2% (14 of 31) showed concurrent ventral spread. However, there was no isolated contrast filling of the ventral epidural space. In addition, when contrast patterns in the L4-5 and L5-S1 levels were separately analyzed, a

Table 1. Demographics of the subjects (n=31)

Parameter	Value
Sex	
Male	12
Female	19
Age (yr)	63.3±14.6 (30-85)
Injection level	
L4-5	20
L5-S1	11
Injection side	
Right	13
Left	18
Lesion type	
Lateral HD	17
Central HD	14
VAS at baseline	6.23±1.33 (4-9)

Values are presented as number or mean±standard deviation (range).

HD, herniated disc; VAS, visual analog scale.

significantly greater tendency was observed for the contrast to expand ventrally at the L5-S1 (8 of 11) level than at the L4-5 (6 of 20) level, according to the results of Fisher exact test (p=0.031, OR=6.2).

In the analysis of efficacy of SP TFESI, whether or not ventral expansion of contrast was present, the changes in the VAS scores and the proportion of pain reduction at 2 weeks after SP TFESI were not statistically different (p>0.1) (Table 2).

In the vertical contrast spread analysis, at L4-5 level, the mean grade of contrast spread was 1.40 ± 1.05 in a cephalic direction and 1.55 ± 1.32 in a caudal direction. At L5-S1 level, the mean grade was 1.73 ± 1.19 and 1.64 ± 0.92 in the cephalic and caudal directions, respectively. In an analysis of the correlation between the vertical spread pattern and treatment effect according to the dorsoventral spread (dorsal isolated and ventral expansion), the grades for cephalic extension in the ventral expansion group showed a meaningful positive correlation with proportional pain reduction (r=0.676, p<0.01) (Table 3).

The changes in the VAS scores along with the degree of vertical contrast spread on each direction is summarized, and listed in Table 4.

In addition, the location of HD (central or lateral HD) was not found to affect the anteroposterior spread of contrast in the epidural space at any level (Fisher exact test, both p>0.1 for L4-5 and L5-S1).

In the extent of vertical spread, the mean grades for

		VAS											
Injection level	Contrast pattern	Dro	After 2 weeks	Reduction									
		Ple	Aller 2 weeks	Δ	Rate (%)								
L4-5	Ventral expansion (n=6)	6.3±1.5	3.0±0.9	-3.33	52.8								
	Dorsal isolated (n=14)	5.9 ± 1.4	2.9±1.0	-2.93	49.9								
	p-value	0.505	0.882	0.394	0.619								
L5-S1	Ventral expansion (n=8)	6.6±1.3	3.6±0.9	-3.00	45.4								
	Dorsal isolated (n=3)	6.7±0.6	4.0±1.7	-2.67	40.3								
	p-value	0.960	0.641	0.628	0.596								
Total	Ventral expansion (n=14)	6.5±1.3	3.4 ± 0.9	-3.14	48.6								
	Dorsal isolated (n=17)	6.0±1.3	3.1±1.2	-2.88	48.2								
	p-value	0.307	0.539	0.434	0.941								

Values are presented as mean±standard deviation.

Student t-test was used to compare the values for each contrast pattern group.

SP TFESI, superoposterior transforaminal epidural steroid injection; VAS, visual analog scale; Δ , changes in VAS value from before and 2 weeks after TFESI.

cephalic and caudal spreads showed no statistical difference between the central and lateral HD, when analyzed according to each level and direction of spread (Student t-test).

No major complications were reported.

DISCUSSION

Recently, several studies reported that SP TFESI is a safer alternative to the conventional SA approach, because of the lower risk of vascular complications [20]. This study aimed to investigate the contrast spreading pattern and its correlation with efficacy for SP TFESI. Contrast was examined for the spread to both the ventral and dorsal epidural spaces, and in cephalic and caudal directions. Clinical efficacy was significantly correlated with the cephalic spreading pattern. These results indicate that the spreading of injectate could be one of the factors determining the clinical efficacy in SP TFESI.

Several previous studies reported conflicting results for differences in clinical efficacy between SA and SP TFESI.

Table 3. Pearson correlation analysis for vertical dispersal extent and degree of proportional pain reduction with SP TFESI according to dorsoventral contrast pattern type

Contrast pattern type	Cephalic	Caudal
Dorsal isolated (n=17)	0.434	0.187
p-value	0.082	0.473
Ventral expansion (n=14)	0.676*	-0.370
p-value	0.008	0.192

SP TFESI, superoposterior transforaminal epidural steroid injection. *p<0.01. [5,15]. Desai et al. [15] reported that the efficacy of SA TFESI was superior to that of SP TFESI and suggested that ventral and greater extent of contrast spread would assure better therapeutic efficacy. However, in this study, the ventral spreading pattern was not correlated with the clinical efficacy in SP TFESI. The ventral spreading of contrast was observed in 45.2% of the cases in this study, which is less than the 61.4% in the study by Desai et al. [15]. In addition, ventral expansion was more commonly observed at the L5-S1 level than at the L4-5 level. According to a study on the morphometrics of the lumbar spinal canal, the lower lumbar spinal canal has a greater midsagittal and interpedicular diameter through L3 to L5 level [21]. The spinal canal space at the L5-S1 level is larger than the space at the L4-5 level, and the less constricting epidural space may permit easier ventral expansion of contrast. Since TFESI was performed more frequently at the L4-5 level than at the L5-S1 level, we assumed that ventral spreading patterns were observed in a relatively small number of patients in this study. However, further research is needed to determine the correlation between the differences in the epidural space and contrast spreading patterns.

Several previous studies examined the vertical direction of contrast spread in TFESI. Two studies reported a tendency for more cephalic spread [15,22]. However, Vassiliev [17] and Gajraj [23] reported preferential spread of contrast onto the medially located nerve roots (lower level of nerve root). Furman et al. [19] observed an equivalent extent of spread in both the directions in the presence of 1 mL of contrast. In this study, the contrast medium spread to a similar degree on our grading system in both the cephalic and caudal directions.

The vertical contrast spread pattern analysis showed

Decision of the state of		wear of vAS changes													
D	egree of vertical	L4	-5	L5-	S1	Total									
	spread	Cephalic	Caudal	Cephalic	Caudal	Cephalic	Caudal								
	Grade 0	-2.50 (4)	-2.80 (5)	(0)	-4.00 (1)	-2.50 (4)	-3.00 (6)								
	Grade 1	-2.57 (7)	-3.40 (5)	-2.71 (7)	-2.50(4)	-2.64 (14)	-3.00 (9)								
	Grade 2	-3.57 (7)	-2.71 (7)	-3.00 (2)	-3.25 (4)	-3.44 (9)	-2.91 (11)								
	Grade 3	-4.00(1)	(0)	(0)	-2.50 (2)	-4.00(1)	-2.50 (2)								
	Grade 4	-4.00(1)	-3.67 (3)	-3.50(2)	(0)	-3.67 (3)	-3.67 (3)								

N/ CTUAO 1

Table 4. Mean of VAS changes according to the degree of vertical extent of contrast spreads

The numbers of patients with each degree of vertical spread are in parentheses. VAS, visual analog scale.

a significant correlation between the cephalad spreading and clinical efficacy in the ventral expansion group. As mentioned above, Desai et al. [15] confirmed that the more level of vertebra covered by the contrast would lead to a better clinical effect. More amount of medication passed into the epidural space would eventually lead to increased flow of medication into the adjacent nerve roots as well as the nerve root of the target level. In concordance to this, we assume that the positive correlation between the vertical contrast extent and therapeutic effect is a reasonable phenomenon.

In this study, since most patients who underwent L4-5 TFESI were thought to have L5 radiculopathy, the therapeutic effects might not be the direct results of TFESI on the L4 nerve root. In these cases, more medial spread of medication might lead to a more successful reduction of the L5 root symptoms, because the L5 root is located medial to the L4 root at L4-5 level. According to the study of Vassiliev [17], volume-spread relationships were observed, when L4-5 TFESI was performed. Contrast spread onto the L5 nerve root in 46.1% with 1 mL of contrast and in 61.5% with 2-3 mL of contrast. No spread of contrast was observed onto the L3 nerve root. In this study, though we used 1 mL of contrast for evaluating the contrast spread patterns, 2.5 mL of medication was used for the therapeutic purpose, and such volume would sufficiently affect the L5 nerve root with L4-5 TFESI.

The effect of positional type of HD (central or lateral) was also evaluated on the contrast spread. Prior studies showed the similar results. Botwin et al. [22] reported no significant difference in the overall flow patterns between the patients with lumbar spinal stenosis and HD by using conventional TFESI. Furman et al. [19] found no statistical significance in the contrast volume needed to reach specific landmarks with or without the presence of central or foraminal stenosis. Similar to prior studies, SP TFESI with central and lateral HD showed did not show any statistical differences in the contrast spread in the ventral or dorsal direction or in vertical extent of spread.

This study has the following limitations. First, contrast has a higher viscosity than the medication. This difference can cause unexpected spread patterns in the injected medication. Moreover, we only captured a fluoroscopic image immediately after the completion of SP TFESI. Because both the contrast and medication would spread further over time, we may have under-estimated the extent of dispersion. However, because the lower viscosity of the medication and the time effect would result in at least equivalent or larger dispersion, we think that our interpretation is still valid or can be extended. If extended, based on adequate spreading, the significance of dividing the anterior and posterior epidural space for evaluation of contrast dispersion would lessen and enhance our result, thereby showing no difference in the proportion of pain reduction between the dorsal spreading and ventral expansion. Another limitation is that the pathologic level of radiculopathy was not clear in some patients. No definite correspondence was observed between the electrophysiologic study, MR images, and clinical symptoms. If cases with more definitive levels of radiculopathy are recruited in a larger study, the results would be demonstrated more clearly.

In SP TFESI, achieving ventral contrast spread did not guarantee better treatment effects; however, that the extent of cephalic spread in the ventral expansion group was positively correlated with the proportion of pain reduction.

Since this study analyzed contrast spread patterns, it should provide data for use in further research. In particular, because SP TFESI is mainly used in difficult cases with severely degenerated vertebrae, it would be meaningful to compare the therapeutic efficacy of SP TFESI and SA TFESI in the patients with severe foraminal stenosis in future study.

SUPPLEMENTARY MATERIALS

Supplementary materials can be found via http://doi. org/10.5535/arm.2017.41.3.413. Table S1. Patients' clinical presentations, results of MRI and EMG, and epidural steroid injection level.

CONFLICT OF INTEREST

No potential conflict of interest relevant to this article was reported.

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		Injection level	L4-5	L4-5	L4-5	L5-S1	L4-5	L4-5	L4-5	L4-5	L5-S1	L4-5	L4-5	L4-5	L4-5	L5-S1	L5-S1	L5-S1	L4-5	L4-5	L4-5	L4-5	L4-5	L4-5	L5-S1	L5-S1	L5-S1	L5-S1	L5-S1	L4-5	L4-5	L5-S1	L4-5		al steroid:
		Anatomical difficulty for ESI	ı	I	ı	ı	ı	High iliac crest	I	ı	ı	ı	Large L5 transverse process		High iliac crest	ı	ı	I	ı	I	High iliac crest	ı	ı	I	ı	ı	ı	ı	ı	I	ı	I	Pseudo-arthrosis	L5-iliac crest	normal; ESI, epidu
	S	Ankle plantar flexor, knee flexo	Х	Х	х	Х	Х	Х	Х	Х	Х	Х	X	Х	Х	Х	Х	Х	Х	Х	Х	Х	х	Х	Х	Х	Х	Х	Х	Х	Х	Х	X		opathy; NL,
	otor weakne	Toe extensor, ankle dorsiflexor	Х	х	х	0	х	х	0	Х	х	х	X	х	0	х	х	х	х	х	х	х	x	х	х	х	х	х	х	х	х	х	x		; radi, radicul
	M	Knee extensor, hip adductor	X	х	x	х	x	х	X	Х	х	х	x	x	X	х	х	х	х	х	х	х	x	х	х	х	х	х	х	х	х	х	х		al; Lat, lateral
	ange	Posterior thigh, calf, heel	X	х	х	\bigtriangledown	х	х	х	х	\bigtriangledown	х	х	х	х	х	x	х	х	х	х	x	x	х	х	\triangleleft	х	x	х	х	х	х	\bigtriangledown		t; Cent, centra
pardo arm (oru	or sensory ch	Lateral thigh, lateral lower leg, dorsum of foot	0	0	0	0	0	0	0	0	0	0	0	x	0	0	0	0	0	0	0	0	0	0	0	⊲	0	0	0	0	0	0	0		tt, right; Lt, left
	Pain	Medial lower leg	х	х	х	х	\bigtriangledown	х	х	х	х	\bigtriangledown	x	0	х	х	х	х	х	Х	х	х	\bigtriangledown	х	х	х	Х	х	х	х	х	Х	х		nyography; I
		EMG	NL	L5 radi	NL	L5 radi	NL	NL	L5 radi	NL	NL	NL	NL	L4 radi	L5 radi	NL	L5 radi	NL	NL	NL	NL	L5 radi	NL	L5 radi	L5, S1 radi	NL	NL	NL	NL	L5 radi	NL	NL	L5 radi		EMG, electror
and brocottan		MRI	L4-5, Lat	L4-5, Lat	L4-5, Lat	L5-S1, Lat	L4-5, Lat	L4-5, Cent	L4-5, Lat	L4-5, Lat	L5-S1, Cent	L4-5, Lat	L4-5, Cent	L4-5, Lat	L4-5, Cent	L4-5, Cent	L4-5, Cent	L4-5, Cent	L4-5, Lat	L4-5, Lat	L4-5, Cent	L4-5, Lat	L4-5, Lat	L4-5, Lat	L5-S1, Cent	L5-S1, Cent	L5-S1, Lat	L4-5, Cent	L4-5, Cent	L4-5, Lat	L4-5, Lat	L4-5, Cent	L4-5, Cent		ce imaging; I
		Side	Lt	Rt	Lt	Rt	Lt	Rt	Lt	Lt	Lt	Rt	Rt	Rt	Rt	Lt	Lt	Rt	Lt	Rt	Lt	Rt	Lt	Lt	Lt	Rt	Lt	Lt	Rt	Lt	Rt	Lt	Lt		onan
		Age (yr)	36	85	99	64	69	53	68	99	81	69	69	63	70	59	65	52	33	74	35	74	77	68	73	60	63	50	79	30	79	52	80		ic rest
The second		Sex	М	ц	ц	ц	ц	ц	щ	ц	ц	Σ	Μ	ц	Μ	щ	И	ц	ц	ц	И	Σ	Μ	ц	ц	ц	И	ц	Σ	ц	М	Σ	ц		gneti
		Patient no.	1	7	3	4	5	9	7	8	6	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31		MRI, ma

Table S1. Patients' clinical presentations, results of MRI and EMG, and epidural steroid injection level

þ injection; o, present; x, not present; Δ , equivocal or not always.