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Clinical outcome of percutaneous endoscopic lumbar decompression in treatment of elderly patients with lumbar spinal stenosis: a matched retrospective study

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Abstract

Purpose This retrospective cohort study aimed to evaluate the efficacy and safety of percutaneous endoscopic lumbar decompression (PELD) in elderly patients with lumbar spinal stenosis (LSS).

Study Design A matched retrospective study.

Setting The research was conducted in Beijing Chaoyang Hospital, Capital Medical University, China.

Methods This study included patients treated with PELD for LSS from September 2016 to September 2020. Patients with LSS aged \geq 80 years were screened according to the inclusion and exclusion criteria as the study group, and then the same number of patients with LSS aged 50-80 years were matched according to gender, stenosis type, and surgical segment as the control group. Preoperative patient status was assessed using the Charlson comorbidity index (CCI) and the American Society of Anesthesiologists (ASA) physical status classification score. Clinical outcomes were assessed using the visual analog scale (VAS), Oswestry Disability Index (ODI) scores, modified Macnab criteria, radiological parameters and complication rates. Results A total of 624 LSS patients met the screening criteria between September 2016 and September 2020, with 47 LSS patients \geq 80 years old serving as the study group. Forty-seven LSS patients aged 50–80 years were matched to the study group according to gender, stenosis type, and stenosis segment. The CCI score (1.77 ± 1.67) and ASA classification (2.62 ± 0.74) of the study group were significantly higher than the CCI score (0.66 ± 0.96) and ASA classification (1.28 ± 0.54) of the control group, and the difference was statistically significant. Compared with preoperative data, postoperative ODI, leg pain VAS scores and back pain VAS scores were significantly improved in both groups (p < 0.05). However, no significant difference was found between two groups in preoperative and postoperative ODI, leg pain VAS scores and back pain VAS scores (p > 0.05). The operation time and postoperative hospital stay in control group were significantly lower than those in study (p < 0.05), but there was no significant difference in blood loss between the two groups (p > 0.05). Besides, overall radiological parameters were comparable in elder and younger patients (p > 0.05), and disc height (DH), lumbar lordosis and segmental lordosis decreased after two year follow-up in both groups (p < 0.05). In addition, complication rates were similar between the two groups (p > 0.05), and no serious complications and deaths were found.

Limitations Single-centre retrospective design, non-randomized sample, small sample size.

Conclusion Although elderly LSS patients (\geq 80 years old) are less fit and have more comorbidities, satisfactory outcomes can be achieved with PELD, comparable to those of LSS patients < 80 years old, and without increased complications.

Keywords Lumbar spinal stenosis · Percutaneous endoscopic lumbar decompression · Elderly patients · Clinical efficacy

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Introduction

Population aging is the inevitable result of population transformation, and it is an important issue facing human society in the twenty-first century. With the development of technology, people over the age of 80 are the fastest growing age group worldwide, which leads to an increase in the incidence of age-related diseases. Lumbar spinal stenosis (LSS) is a

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common age-related degenerative disease [1]. In the national low back pain study, 22% of patients with chronic low back pain were ultimately diagnosed with LSS [2]. LSS can cause moderate to severe pain that affects the quality of life of patients [3]. Elderly patients with LSS usually suffer from long-term pain and underlying diseases, and conservative treatment results are poor [4]. Patients with LSS who fail conservative treatment often require surgery.

Open decompression (OD) has been considered the gold standard surgical option for LSS [5]. OD can relieve nerve damage and improve associated pain by relieving lumbar spinal stenosis and removing excess tissue [6]. However, due to the disadvantages of OD, such as large trauma, long operation time, high blood loss, and increasing the incidence of complications in elderly patients, most elderly LSS patients cannot achieve satisfactory results after OD treatment [7-9]. Therefore, it is still necessary to explore new surgical solutions for elderly LSS patients with many comorbidities. With the improvement of surgical techniques, optical equipment and surgical instruments, minimally invasive techniques for the spine have developed rapidly. The percutaneous endoscopic lumbar decompression (PELD) method has become a popular minimally invasive technique for the treatment of lumbar degenerative diseases [10, 11]. According to studies, PELD has the advantages of less tissue trauma and shorter operation time than OD [12, 13]. Additionally, studies have found that PELD and OD have similar efficacy in the treatment of LSS, but less complications and mortality than OD [14]. PELD has emerged as an important alternative treatment for LSS. However, no relevant studies have reported the efficacy and safety of PELD in the treatment of elderly LSS patients (especially those with multiple comorbidities and poor physical conditions). Therefore, we designed a gender, stenosis type and surgical segment-matched retrospective study to investigate the clinical outcomes of elderly LSS patients (\geq 80 years) treated with PELD in order to provide clinical guidance for surgeons in the treatment of elderly LSS patients.

Patients and methods

Patients

A retrospective analysis was performed on patients with LSS who received PELD from September 2016 to September 2020. The inclusion criteria for this study were: 1. Patients aged \geq 50 years; 2. Clinically diagnosed with symptomatic LSS and ineffective conservative treatment for three months; 3. Patients receiving PELD therapy. Patients with cervical and thoracic spinal diseases; history of previous spinal surgery; spinal deformity, spondylolisthesis and lumbar instability; combined lumbar infection and tumour;

and patients with incomplete data or lost to follow-up were excluded. During this period, a total of 624 patients with LSS underwent PELD surgery, including 47 patients with LSS \geq 80 years as the study group. Then, 47 LSS patients aged 50–80 years who matched the study group in terms of gender, stenosis type, and stenosis segment were selected as the control group. All patients were followed up for two years after surgery. This study was approved by the ethics committee, and all patients signed informed consent.

Surgical technique

PELD procedures were all performed by a senior spine surgeon (ZL) with sufficient experience, at a single level, under local anaesthesia, with the patient placed in a prone position. The patient can communicate with the surgeon during the entire operation to prevent intraoperative nerve root injury. For patients with multilevel stenosis, to determine the level most responsible to symptoms, best suiting for the procedure, a transforaminal selective nerve root block was performed.

The entry point was set at 12 to 14 cm lateral to spinal middle line at the index intervertebral level. Then puncture needle was inserted into the superior articular process (SAP) of the targeted segment. Through the puncture needle, serial cannulated dilators were inserted into the SAP. The ligamentum flavum and the ventral elements on the SAP were removed. Then, a tubular retractor with an outer diameter of 7.9 mm was passed over the dilators and secured to the upper lamina and burrs were used to further enlarge the foramen if necessary. The herniated disc, parts of posterior longitudinal ligament, posterior upper margin of the inferior vertebral body, posterior lower margin of the superior vertebral body (if necessary) and the dorsal ligamentum flavum were removed with rongeur, endoscopic forceps, endoscopic bone knife or a highspeed drill. Finally, the entire nerve root and dural sac was probed to ensure complete decompression. Adequate irrigation and haemostasis were performed to reduce postoperative infection and hematoma.

Clinical evaluation

Demographic data of all study participants were collected, including gender, age, Obesity (BMI \geq 28), smoking, length of postoperative hospital stays, lesion segment, operative time, blood loss, stenosis type, Charlson comorbidity index (CCI) [15], the American Society of Anesthesiologists (ASA) physical status classification score [16]. The visual analog scale (VAS) was used to assess leg pain and low back pain. The Oswestry Disability Index (ODI) was used to assess clinical functional status. The surgical satisfaction was assessed using the modified Macnab criteria. In addition, postoperative complications were also assessed in this study.

Radiological evaluation

Disc degeneration status were identified by disc height (DH) and the Pfirrmann grade of disc. DH was measured as the mean distance of the anterior and posterior disc heights on mid-sagittal MRI T2-weighted sequence. The Pfirrmann grading system ranging from I to V was also used in assessing disc degeneration [17]. Lumbar lordosis was measured as the angle between superior endplate of L1 and the superior endplate of S1, and segmental lordosis was measured as the angle between the inferior endplate of the superior vertebrae and the superior endplate of the inferior vertebrae at the index level by lateral radiographs. Newly developed disc herniation, lumber stenosis, spondylolisthesis, and lumbar instability were detected by the comparison of the final follow-up and preoperative MRI, lateral radiographs, or flexion–extension radiographs.

Statistical analysis

Data analysis was performed with GraphPad Prism 8 (GraphPad Software, San Diego, CA). Data are presented as mean \pm standard deviation (SD). Data comparisons between preoperative and postoperative time points were analyzed using paired samples t-tests or McNemar-Bowker tests. Comparisons between the two groups were assessed using an independent samples t-test. The chi-square test was used to compare discontinuous variables between the two groups. The p < 0.05 indicated that the difference was statistically significant.

203

Results

A total of 624 LSS patients met the screening criteria between September 2016 and September 2020, with 47 LSS patients \geq 80 years old serving as the study group. Forty-seven LSS patients aged 50–80 years were matched to the study group according to gender, stenosis type, and stenosis segment.

Those patient baseline characteristics were shown in Table 1. The mean age of patients in study and control group was 83.26 ± 2.53 and 64.55 ± 8.32 years, respectively. The most common lumbar spine level in study and control group was L4-5 (study group: n = 36, control group: n = 36), followed by L5-S1 (study group: n = 6, control group: n = 6) and L3-4 (study group: n = 5, control group: n = 5). All patients in study group had comorbidities, including hypertension, diabetes mellitus, cardiovascular diseases, cerebrovascular disease, pulmonary disease, liver / gastrointestinal diseases, renal insufficiency, history of anticoagulants or antiaggregants use, osteoporosis and cancer history (Table 2). In control group, 29 patients (61.7%) had comorbidities. The incidence of comorbidities was significantly different between the two groups (P < 0.05). The mean CCI score and ASA classification in study group were 1.77 ± 1.67 and 2.62 ± 0.74 , respectively; while the mean CCI score and ASA classification in control group were 0.66 ± 0.96 and 1.28 ± 0.54 , respectively. There were statistically significant differences in CCI score and ASA classification between the two groups (p < 0.05). The operative time and postoperative

Table 1Baseline characteristicsof patients in both groups

	Study group	Control group	p-Value	
Age (years)	83.26 ± 2.53	64.55 ± 8.32	< 0.0001*	
Sex (male/ female)	22/25	22/25	Matched	
Obesity (BMI \geq 28)	3 (6.4%)	11 (23.4%)	0.0205*	
Smoking, n	4 (8.5%)	13 (27.7%)	0.0159*	
Duration of symptoms (months)	27.26 ± 27.85	15.77 ± 12.9	0.0119*	
Type of stenosis			Matched	
Central stenosis	26 (55.3%)	26 (55.3%)		
Lateral recess stenosis	21 (44.7%)	21 (44.7%)		
Operative level			Matched	
L3-4	5 (10.6%)	5 (10.6%)		
L4-5	36 (76.6%)	36 (76.6%)		
L5-S1	6 (12.8%)	6 (12.8%)		
Blood loss (ml)	7.77 ± 5.19	9.79 ± 5.51	0.0705	
Operative time (minutes)	147.8 ± 48.05	123.1 ± 34.05	0.0050*	
ASA classification	2.62 ± 0.74	1.28 ± 0.54	< 0.0001*	
CCI	1.77 ± 1.67	0.66 ± 0.96	0.0002*	
Postoperative hospital stay (day)	8.57 ± 7.06	4.38 ± 1.73	0.0002*	

BMI: body mass index; ASA: American Society of Anesthesiologists; CCI: Charlson comorbidity index

* Statistical significance (*P* value < 0.05)

Table 2 Comorbidities in the two groups of patients

Comorbidities	Study group	Control group	<i>p</i> -Value	
Hypertension	38 (80.9%)	7 (14.9%)	< 0.0001*	
Diabetes mellitus	14 (29.8%)	10 (21.3%)	0.4785	
Cardiovascular disease	25 (53.2%)	4 (8.5%)	< 0.0001*	
Cerebrovascular disease	14 (29.8%)	4 (8.5%)	0.0166*	
Pulmonary disease	7 (14.9%)	3 (6.4%)	0.3161	
Liver / Gastrointestinal	6 (12.8%)	2 (4.3%)	0.2673	
Renal insufficiency	5 (10.6%)	2 (4.3%)	0.4349	
Urinary tract infection	5 (10.6%)	1 (2.1%)	0.2035	
Peripheral vascular disease	5 (10.6%)	4 (8.5%)	> 0.9999	
Use of anticoagulants or antiaggregants	12 (25.5%)	2 (4.3%)	0.0073*	
Osteoporosis	23 (48.9%)	10 (21.3%)	0.0094*	
Any malignancy	3 (6.4%)	1 (2.1%)	0.6168	
Total	47 (100%)	25 (53.2%)	< 0.0001*	

* Statistical significance (*P* value < 0.05)

hospital stays of the control group were significantly lower than those of the study group (p < 0.05). In addition, there was no significant difference in blood loss between the two groups (p > 0.05).

There were no major complications or procedurerelated deaths in either group. Eight patients in study group had general complications including cardiovascular disease, pulmonary disease, liver/gastrointestinal disease, renal insufficiency, urinary tract infection and deep venous thrombosis; four patients in control group had general complications, and there was no statistical difference between the two groups (p > 0.05, Table 3). In addition, surgical complications occurred in 11 patients (7 patients (14.9%) in study group, four patients (8.5%) in control group, p > 0.05, Table 3). Six patients from both groups underwent reoperation for different reasons: incomplete decompression (2 patients), contralateral symptoms after decompression (1 patient) and recurrent disc herniation (3 patients).

Compared with preoperative data, both groups showed significant improvements in ODI, leg pain VAS scores and back pain VAS scores at three months, one year and two years postoperatively (p < 0.05, Fig. 1). ODI, leg pain VAS scores and back pain VAS scores in study group decreased from preoperative 67.94 ± 5.57 , 6.17 ± 1.03 and 6.15 ± 0.86 to two years postoperative 21.17 ± 4.33 , 2.26 ± 1.01 and 2.3 ± 1.16 , respectively. In control group, ODI, leg pain VAS scores and back preoperative 69.51 ± 5.83 , 5.98 ± 1.01 and 6.04 ± 0.83 to postoperative 20.55 ± 4.89 ,

Table 3 Complications in the two groups of patients

Complications	Study group	Control group	p-Value	
General complications	8 (17.0%)	4 (8.5%)	0.22	
Cardiovascular disease	2	0		
Pulmonary disease	2	0		
Liver / Gastrointestinal	1	1		
Renal insufficiency	2	0		
Urinary tract infection	2	2		
Deep venous thrombosis	1	1		
Surgery-related complica- tions	7 (14.9%)	4 (8.5%)	0.34	
Dural tear	2	2		
Hematoma	1	0		
Neurologic deficit	1	1		
Infection	1	0		
Reoperation	4	2		

 2.11 ± 0.91 and 2.17 ± 0.99 , respectively. No significant differences were found in ODI, leg pain VAS score and back pain VAS score between the two groups in the preoperative, three month, one year and two year postoperative periods (p > 0.05, Fig. 1).

According to the modified MacNab criteria, 39 patients (83%) in study group had a good-to-excellent outcome and 42 patients (89.4%) in control group had a good-to-excellent outcome, with no statistical difference between the two groups (p > 0.05, Fig. 2). Representative case is presented in Fig. 3.

Radiological parameters were further evaluated (Table 4). Compared to preoperative disc degeneration status, DH at final follow-up decreased significantly (study group: $[7.6 \pm 2.3]$ mm vs. $[7.9 \pm 2.2]$ mm, P < 0.01; control group: $[8.2 \pm 1.6]$ mm vs. $[8.5 \pm 1.6]$ mm, P < 0.01). Although more patients suffered from Pfirrmann grading \geq IV at the index disc in study group compared with control group, no significant differences were identified in the two groups between preoperative and final follow-up data. Both lumbar lordosis and segmental lordosis were significantly decreased at final follow-up in the study and control group, yet there was no difference between the two groups. There were five patients (3 in study group and 2 in control group) suffered from newly developed disc herniation, and four patients (2 in study group and 2 in control group) suffered from newly developed lumbar stenosis. Furthermore, four patients developed spondylolisthesis (1 in study group and 3 in control group), and one patient in study group developed lumbar instability postoperatively. None of disc herniation, lumbar stenosis, spondylolisthesis, and lumbar instability significantly differed between the two groups.

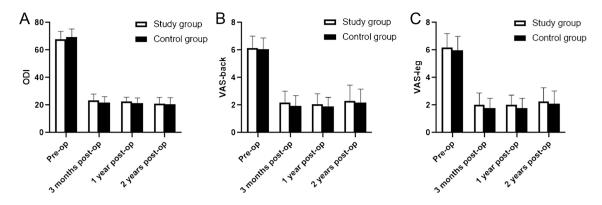
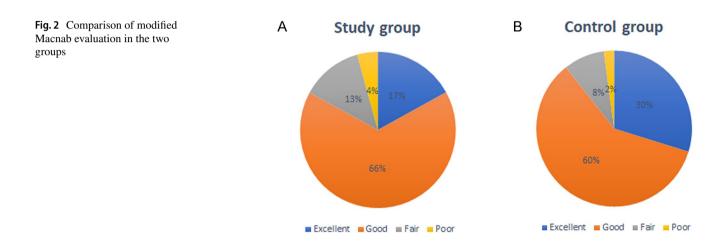


Fig. 1 Preoperative and postoperative clinical outcomes: A. ODI, B. back pain VAS score, C. leg pain VAS scores. VAS, visual analogue scale; ODI, Oswestry disability score



Discussion

Few studies have reported the efficacy and safety of PELD in the treatment of elderly LSS. Therefore, this study is the first to compare the efficacy and safety of PELD in treating LSS patients ≥ 80 years old and < 80 years old. Our results showed that LSS patients aged ≥ 80 years can achieve satisfactory efficacy after PELD treatment, which is comparable to that of LSS patients aged < 80 years without increased complications.

OD is considered the gold standard for surgical treatment of LSS in patients with LSS who are refractory to conservative treatment or in severe disease [18]. Many studies have demonstrated that OD is effective in the treatment of LSS [19, 20]. However, laminectomy due to OD may affect the stability of the spine, resulting in varying degrees of injury and complications [21]. In addition, because elderly LSS patients are often accompanied by poor physical conditions and many comorbidities, the large trauma and blood loss of OD surgery will be detrimental to the recovery of elderly LSS patients [22, 23]. With the development of technology and concepts, PELD has become a treatment for degenerative diseases of the lumbar spine and has been shown to be effective for LSS [24, 25]. PELD is becoming more and more popular in LSS due to the advantages of short operative time, low surgical risk, minimal impact on spinal stability, quick postoperative recovery, and fewer complications [26, 27]. Lv et al. compared the outcomes of PELD versus fenestration for the treatment of external lumbar indentation stenosis in elderly patients over 75 years of age [14]. They found that both PELD and fenestration showed good clinical outcomes for the treatment of lumbar scoliotic stenosis, with fewer complications from PELD. A retrospective study showed that the clinical outcomes of PELD for external pit stenosis were excellent, with an improvement in the VAS total score from 7.9 ± 1.2 to 2.3 ± 1.0 and an ODI decrease from 69.1 ± 7.3 to 24.7 ± 6.4 [28]. However, there are few studies on the efficacy and safety of PELD in the treatment of elderly LSS. Only one study reported the efficacy of PELD in patients over 75 years of age. They found that PELD was safe and effective for LSS treatment

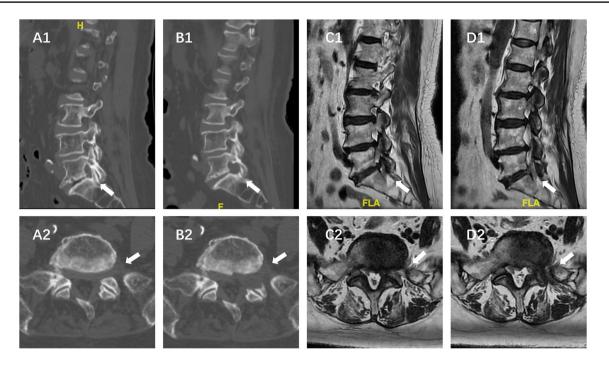


Fig. 3 Percutaneous endoscopic lumbar decompression performed on a 81-year-old female patient diagnosed with L5/S1 left lateral recess stenosis. A1, preoperative CT on sagittal scans. A2, preoperative CT on axial scans. B1, postoperative CT on sagittal scans. B2, postopera-

tive CT on axial scans. C1, preoperative MRI on sagittal scans. C2, preoperative MRI on axial scans. D1, postoperative MRI on sagittal scans. D2, postoperative MRI on axial scans

Radiological parameters	Study group $(N=47)$			Control group $(N=47)$		
	Pre-op	Final follow-up	Р	Pre-op	Final follow-up	Р
Disc degeneration						
Disc height, mm	7.9 ± 2.2	7.6 ± 2.3	< 0.01	8.5 ± 1.6	8.2 ± 1.6	< 0.01
Pfirrmann grading≥IV, n (%)	42 (89.4%)*	46 (97.9%)*	0.13	34 (72.3%)	37 (78.7%)	0.25
Lumbar lordosis, °	34.5 ± 14.9	32.3 ± 13.9	< 0.01	33.6 ± 14.2	32.7 ± 13.6	< 0.01
Segmental lordosis, °	9.4 ± 4.8	$8.3 \pm 5.1^{*}$	< 0.01	11.1 ± 4.8	10.9 ± 5.0	< 0.01
Newly disc herniation, n (%)	-	3 (6.4%)	-	-	2 (4.3%)	-
Newly lumber stenosis, n (%)	-	2 (4.3%)	-	-	2 (4.3%)	-
Newly spondylolisthesis, n (%)	-	1 (2.1%)	-	-	3 (6.4%)	-
Newly lumbar instability, n (%)	-	1 (2.1%)	-	-	0 (0)	-

* indicates statistical difference (P < 0.05) compared to control group

Pre-op pre-operative, Post-op post-operative

in elderly patients, with significantly lower postoperative VAS scores for both low back and leg pain compared to preoperatively [25]. Similar to their results, our study also found that PELD has good clinical efficacy in elderly LSS patients (\geq 80 years). Patients with LSS \geq 80 years had significantly improved postoperative ODI, leg pain VAS scores, and back pain VAS scores after PELD treatment compared with preoperative data. According to Macnab criteria, 39 patients (83%) had a good-to-excellent outcome. Moreover, Li et al. also showed that elderly LSS

patients can achieve the same satisfactory outcomes as younger patients after PELD treatment [25], which is similar to the results of our study. Our results also show that LSS patients ≥ 80 years old can achieve the same curative effect as patients < 80 years old after PELD treatment, and there is no significant difference in complications between the two groups, indicating that PELD treatment of LSS patients ≥ 80 years old is safe and effective.

Patients \geq 80 years old had more comorbidities, and the ASA grade and CCI index were higher than younger patients.

It has been reported that higher CCI and ASA scores are associated with higher likelihood of complications [29]. Studies have demonstrated that the risk of postoperative severe complication is independently associated with heart disease and anticoagulant use [30, 31]. A nationwide populationbased study showed that after LSS surgery, patients with diabetes were approximately 1.35 times more likely to die than those without diabetes [32]. In our study, all patients in study (\geq 80 years) had comorbidities, and the CCI and ASA scores were higher than those in control group (< 80 years), the difference was statistically significant. However, elderly LSS patients in study could achieve similar efficacy to control group after PELD treatment, and there was no significant increase in complications. These results suggest that PELD can be safely treated in elderly patients with multiple comorbidities without increasing complications.

Complication rate is an important factor in evaluating the safety of surgery. Studies have reported complication rates of 6 to 52% for all LSS decompression procedures and 10 to 20% for patients \geq 80 years [20, 33]. Previous studies have shown that patients aged ≥ 85 years had more postoperative complications after decompression surgery than other age groups [33, 34]. In our study, we found that the complications of PELD in the treatment of LSS patients \geq 80 years old are rare, the incidence of general complications is 17.0%, and that of surgery-related complications is 14.9%. Moreover, we also found that there was no significant difference in the risk of complications between patients \geq 80 years old and patients < 80 years old. In our study, for general complications, we keep close contact with physicians and actively treat related complications. For the surgery-related complications, different types adopt different ways to deal with them. Haematoma is a rare complication of spinal surgery and should be detected promptly and surgically removed as soon as possible. In our study, one patient developed a haematoma, numbness and weakness of the lower extremities and decreased muscle strength after PELD. Surgery was performed urgently to remove the haematoma and after that the symptoms of nerve root compression were relived. To reduce the risk of postoperative bleeding and hematoma formation, we usually decrease the irrigation water pressure to detect and haemostasis potential bleeding before the end of the operation. Dural tear is a common complication associated with surgery. In this study, there were four patients with dural tears, all of which occurred at the nerve root sleeve and were caused by the severe adhesion. They were treated conservatively by abdominal compression bandaged bed rest for one week, no cerebrospinal fluid leakage and wound infection. For patients who need resurgery, PELD can be used again for revision surgery.

Few studies have evaluated the morphometric effects after PELD. Li et al. found no significant difference of DH, lumbar lordosis and segmental lordosis after PELD during a at least one year follow-up compared to those preoperatively [35]. However, the follow-up period of the study was relatively shorter. In our study, we found all of DH, lumbar lordosis and segmental lordosis were decreased after PELD with a at least two year follow-up. However, no statistical differences were identified between elder and younger patients. We believe this may attribute to the natural degeneration of patients instead of the influence of surgical intervention, and PELD can achieve sufficient morphometric effect in both elder and younger patients with LSS.

There are some limitations in this study. First, this study is a single-centre, small sample size retrospective study and has a short follow-up period. Second, although the two matched groups of individuals had the same gender, stenosis type, and surgical segment, some variables such as the degree of spinal canal and foraminal stenosis could not be controlled, which may have influenced the results. Finally, this study did not evaluate the clinical outcomes of PELD versus OD in the treatment of elderly LSS patients. Therefore, additional studies with larger sample sizes are needed for supplementation and validation.

Conclusion

This study demonstrated that PELD is effective and safe for LSS patients aged ≥ 80 years. In addition, this study also found that PELD has similar benefits and risks in LSS patients over and under 80 years of age. Our results may provide a good treatment option for surgeons treating elderly LSS patients (≥ 80 years) with comorbidities.

Authors' contributions Conceptualization: Lei Wang and Lei Zang; methodology: Lei Wang and Tianyi Wang; formal analysis and investigation: Ning Fan and Shuo Yuan; writing—original draft preparation: Lei Wang and Peng Du; writing—review and editing: Peng Du and Fangda Si; resources: Lei Zang; supervision: Lei Zang.

Data availability The datasets generated during and/or analysed during the current study are available from the corresponding author on reasonable request.

Code availability Not applicable.

Declarations

Ethics approval The research conducted has been performed in accordance with the Declaration of Helsinki. Approval for the study was obtained from the ethics committees of the Beijing Chaoyang Hospital (2021-KE-478). Informed consent to this study was waived because of this was a retrospective study, which was also approved by the institutional ethical review board.

Consent to participate Not applicable.

Consent to publish Not applicable.

Conflicts of interest All authors declare that they have no conflict of interest.

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209

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