Clinical effect evaluation and correlation between preoperative imaging parameters and clinical effect of endoscopic Transforaminal decompression for lumbar spinal stenosis

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Abstract

Background: The objective of this study was to evaluate the clinical effect and correlation between preoperative imaging parameters and the clinical effect of endoscopic transforaminal decompression for lumbar spinal stenosis.

Methods: In this prospective study, 87 patients from Shanxi Province People’s Hospital met the criteria for lumbar spinal stenosis and were recruited from June 2014 to January 2016. These patients underwent endoscopic transforaminal decompression. The clinical symptoms were evaluated by VAS, ODI, and claudication at 3 and 6 months after surgery. The overall clinical efficacy was evaluated using the MacNab score. Yellow ligament thickness and area of the dural sac were examined by MRI. Bony vertebral canal area, real spinal canal area, nerve root canal bony area, nerve root canal real area, distance between the articular joints, and vertebral canal sagittal diameter were examined by CT. The soft tissue invasion ratio of the vertebral canal and the invasion ratio of the nerve root canal were calculated. Correlations between imaging parameters and age, sex, and clinical efficacy were examined.

Results: The MacNab scores were excellent in 47% of cases, good in 34%, generally good in 8%, and poor in 11%. VAS, ODI, and claudication were significantly improved compared with the preoperative values (P<0.01). A significant difference was observed between the 71–81 year age group and the other age groups (P<0.05). There were good correlations between clinical efficacy and vertebral canal sagittal diameter, distance between the articular joints, soft tissue invasion ratio of the vertebral canal, and invasion ratio of the nerve root canal.

Conclusion: Treatment of lumbar spinal stenosis by endoscopic transforaminal decompression can achieve good clinical results. This operation is less effective in patients older than 71 years of age. There were positive correlations between clinical efficacy and the vertebral canal sagittal diameter, the articular joints, soft tissue invasion ratio of the vertebral canal, and invasion ratio of the nerve root canal.

Keywords: Endoscopic transforaminal discectomy, Lumbar spinal stenosis, Imaging parameters
Background

Percutaneous endoscopic lumbar discectomy (PELD) can be used for the treatment of various types of lumbar intervertebral disc herniation. With the improvements of instruments and equipment, percutaneous spinal endoscopic technology can now be used to treat lumbar spinal stenosis, and satisfactory clinical effects are achieved. The advantages of PELD include its minimal invasiveness (8 mm incision), ability to penetrate physiological channels (intervertebral bore), and minimal tissue destruction. Nevertheless, the treatment of lumbar spinal stenosis with PELD is still being questioned by traditional open surgeons. We currently use percutaneous spinal endoscopic technology to treat lumbar spinal stenosis in some patients, and from our preliminary clinical observations, most postoperative patients achieve satisfactory clinical effects, but the actual clinical curative effects still have to be assessed. Lumbar spinal stenosis is caused by bony stenosis (articular process hyperplasia of cohesion, hyperplasia of vertebral rear osteophyte, and calcified discs) and soft tissue factors (yellow ligament proliferous hypertrophy and uncalcified discs). Therefore, we speculate that the reason for poor efficacy may be that intervertebral endoscopic surgery is better for the relief of soft tissue compression and limited for the treatment of osseous compression. Therefore, the objective of this study was to evaluate the clinical effectiveness and correlation between preoperative imaging parameters and the clinical effect of PELD for lumbar spinal stenosis.

Methods

Study design and participants

This was a prospective study of 87 patients with a first diagnosis of lumbar spine stenosis at Shanxi Province People’s Hospital between June 2014 and January 2016.

The inclusion criteria were: 1) 25–85 years of age; 2) unilateral or bilateral symptoms of nerve root with (or without) cauda equina compression; 3) single or double segment stenosis symptoms (patients with three or more stenosed segments undergo open surgery were excluded); 4) intermittent neurogenic claudication; and 5) imaging findings consistent with clinical symptoms in terms of pain location or affected root nerve: i) lateral crypt and/or intervertebral pore stenosis; ii) central vertebral canal and/or mixed spinal stenosis; and iii) failure to 3–6 months of conservative treatment.

The exclusion criteria were: 1) intervertebral instability (in accordance with the White criteria, i.e., over-extension and over-flexion X-ray images in the standing position show that the horizontal displacement of adjacent vertebrae is ≥3 mm or the angle change is ≥15° [1]), osseous infection, mental abnormalities, tumor(s), or communication difficulties; or 2) symptoms of stenosis caused by pure intervertebral disc herniation.

The following rules were applied in the presence of stenosis in more than one segment. 1) If CT and MRI showed two-segment stenosis, and if the clinical symptoms and signs (according to the sensory manifestations and mobility in the innervated areas of the corresponding nerve roots) also met the clinical manifestations of two-segment stenosis, the patients were considered as having two-segment stenosis; they were then included. 2) If CT and MRI showed two-segment stenosis, the clinical symptoms and signs did not show any evident manifestations in the innervated areas of the corresponding nerve roots, and the patient only had the symptom of intermittent neurogenic claudication, but without being able to determine from which segment this symptom was from, the patients were also considered as being two-segment stenosis; they were then included. 3) If CT and MRI showed three or more segment stenosis, but the clinical symptoms and signs showed that they were caused by two-segment stenosis (meaning that the symptoms were restricted to the innervated areas of two segments), the patients were considered as having two-segment stenosis. They were included in the study. 4) If CT and MRI showed three or more segment stenosis, and the clinical symptoms and signs showed that they were caused by three-segment stenosis (meaning that the symptoms were determined as coming from three segments), the patients were considered as having three or more segment stenosis and were excluded. 5) If CT and MRI showed three or more segment stenosis, but the clinical symptoms and signs could not identify the responsible segments, and none of the segments could be ruled out for inducing the disease, the patients were considered with three or more segmental stenosis, and were excluded from the study.

Surgical methods

According to the patient’s preoperative CT and MRI parameters, and their clinical symptoms, the segment of interest for decompression was identified. A physician with experience in treating lumbar spinal stenosis by PELD performed the surgery routinely and ensured complete decompression during the operation. The operation was performed through the intervertebral foramen (Additional file 1: Figure S1).

Preoperative and postoperative functional assessment

Clinical symptoms were assessed using the visual analog pain score (VAS) [2, 3], Oswestry Disability Index (ODI) [4, 5], and claudication distance (m). These factors were evaluated independently by two orthopedic surgeons during the preoperative period and at 3 and 6 months postoperatively. The mean value was used as the standard for evaluating preoperative and postoperative clinical symptoms. The overall clinical efficacy was assessed.
using the MacNab scores [6]. The minimal clinically important difference for ODI is 12.4 [7].

Preoperative imaging parameter measurement
Patients were evaluated by MRI (Germany Siemens 3.0 T superconducting magnetic resonance instrument) and CT (Germany’s Siemens 64 layer spiral CT machine). The following indicators were measured by two radiologists who were blinded to the patient’s condition. CT and MRI analysis software was used to calculate the mean values. MRI was used to measure the yellow ligament thickness and area of the dural sac (Additional file 1: Figure S2). CT was used to measure the bony vertebral canal area, real spinal canal area, nerve root canal bony area, nerve root canal real area, distance between the articular joints, and vertebral canal sagittal diameter (Additional file 1: Figure S3).

Definitions
The soft tissue invasion ratio of the vertebral canal and invasion ratio of the nerve root canal were calculated as: the soft tissue invasion ratio of the vertebral canal = (bony vertebral canal area - real spinal canal area)/bony vertebral canal area; the invasion ratio of the nerve root canal = (nerve root canal bony area - nerve root canal real area)/nerve root canal bony area.

Statistical analysis
SPSS 22.0 (IBM, Armonk, NY, USA) and GraphPad Prism 6 (GraphPad Software, San Diego CA, USA) were used for statistical analysis. Normally distributed data (according to the Kolmogorov-Smirnov test) are presented as means ± standard deviation, while non-normally distributed data are presented as median (IQR). Categorical variables are presented as frequencies. The Friedman analysis was used to compare efficacy indicators of different time points among multiple groups, and Dunn’s test was used for comparison between groups. The comparisons of sex and age groups were performed using two-way repeated measure ANOVA with the Bonferroni post hoc test. Correlation between sex, age, and CT/MRI parameters and correlation between CT/MRI parameters and therapeutic effects were analyzed by Spearman correlation. Multiple linear regression was used to analyze the association among various indicators. Two-sided P-values < 0.05 were considered statistically significant. No multiple testing adjustments were done.

Results
Characteristics of the participants
Of the recruited participants, 45 were male (51.7%), and 42 were female (48.3%). The median age was 58 years (range 43–67 years). Table 1 presents the preoperative characteristics of the participants.

| Table 1 Baseline data and CT/MRI parameters |
|---------------------------------------------|
| Variables                                   | All patients (n = 87) |
| Age (years)                                 | 58 (43,67) |
| Male                                        | 45 (51.7%) |
| Bony vertebral canal area by CT measurement (mm²) | 253.8 ± 52.0 |
| Cross-sectional area of the dural sac by MRI measurement (mm²) | 82.9 (62,5112.5) |
| Nerve root canal bony area by CT measurement (mm²) | 79.0 ± 30.1 |
| Yellow ligament thickness by MRI measurement (mm) | 5.2 (4.3,5.6) |
| Real spinal canal area by CT measurement (mm²) | 35.3 (28.5,45.2) |
| Nerve root canal real area by CT measurement (mm²) | 45.1 (31.3,59.4) |
| Vertebral canal sagittal diameter by CT measurement (mm) | 19.8 (18.1,20.9) |
| Distance between the articular joints by CT measurement (mm) | 16.4 (15.3,18.0) |
| Soft tissue invasion ratio of the vertebral canal | 0.86 (0.82,0.88) |
| Invasion ratio of the nerve root canal | 0.42 ± 0.12 |

Evaluation of surgical efficacy
The MacNab score was excellent in 47% of the patients, good in 34%, generally good in 8%, and poor in 11%. The overall optimal rate was 81.1%.

Postoperative VAS, ODI, and claudication at 3 months were significantly improved compared with the preoperative values (P < 0.01). Postoperative VAS, ODI, and claudication at 6 months were significantly improved compared with the preoperative values (P < 0.01) (Table 2). These results indicate that PELD was effective in these patients. The changes from baseline in ODI scores are greater than the minimal clinical significant change of 12.4.

There was no statistically significant difference between sexes (P > 0.05). On the other hand, in the curative effect comparison among the different age groups, a significant difference was observed between the 71–81 year age group and the other age groups (P < 0.05). The overall analysis revealed that the 71–81-year group had worse postoperative effects than the other age groups (Fig. 1).

Correlation of the preoperative imaging parameters with age, sex, and clinical efficacy
There was no significant correlation between most imaging parameters and sex (P > 0.05). There was a correlation between the soft tissue invasion ratio of the vertebral canal and the invasion ratio of the nerve root canal with sex (P < 0.05) (Table 3).

The actual spinal canal area was positively correlated with age (P < 0.05). The soft tissue invasion ratio of the vertebral canal and the invasion ratio of the nerve root canal was negatively correlated with age (P < 0.05) (Table 4).
A good correlation was observed between VAS and articular joints by CT, soft tissue invasion ratio of the vertebral canal, and invasion ratio of the nerve root canal \((P<0.05)\); ODI and articular joints by CT, soft tissue invasion ratio of the vertebral canal, and invasion ratio of the nerve root canal \((P<0.05)\); and walking distance and articular joints by CT, soft tissue invasion ratio of the vertebral canal, and invasion ratio of the nerve root canal \((P<0.05)\) (Table 5).

**Multiple linear regression of VAS, ODI and walking distance in 6 months**

There was a significant correlation between the soft tissue invasion ratio of the vertebral canal, invasion ratio of the nerve root canal, actual spinal canal area, and nerve root canal real area with VAS \((P<0.05)\) (Table 6).

There was a significant correlation between the distance between the articular joints and the soft tissue invasion ratio of the vertebral canal with ODI \((P<0.01)\) (Table 6).

There was a significant correlation between the distance between the articular joints, soft tissue invasion ratio of the vertebral canal, invasion ratio of the nerve root canal, and real spinal canal area with claudication distance \((P<0.05)\) (Table 6).

These results indicate that imaging parameters are independently associated with the clinical outcomes after PELD for lumbar segment spinal stenosis.

**Discussion**

There is presently no uniform standard regarding the surgical indications for the treatment of lumbar spinal...
stenois, and there are different opinions regarding the treatment of lumbar spinal stenosis with PELD. Polikandriotis et al. [8] examined bleeding, time, pain, and functional improvement before and after PELD compared with conventional surgery, and consider that PELD through the laminae or foramen is safe and effective for the treatment of lumbar spinal stenosis. Polikandriotis et al. [8] microscopically treated lumbar spinal stenosis disease through the intervertebral foramen and noted that the operation time for this type of surgery was short, hemorrhage was minimal, and operative and postoperative complications were slightly more frequent than those with traditional surgery. Thus, these authors concluded that this approach is safe and effective for lumbar spinal stenosis surgery PELD and whether they are the same as for traditional open surgery remain to be defined.

Imaging measurement of lumbar spinal stenosis is an auxiliary and indispensable examination that can be performed for diagnosis. Zhang et al. [12] accurately delineated the boundary of the area and measured the area, angle, and segment length of the lesion using measurement software. Liu [13] measured the bones of 100 corpses, and the diameter of the spinal canal in L4 was 15.31 ± 2.23 mm, and that in L5 was 15.98 ± 2.58 mm. Guo [14] et al. measured the diameter of the L5 vertebral canal to be 16.61 ± 2.42 mm. In the present study, the canal in L4 was 15.44 ± 3.06 mm, and that in L5 was 16.18 ± 2.29 mm. The results showed that measurement of the diameter of the vertebral canal was very similar to that of previous studies, indicating that the CT images are close to the physical specimen measurement.

We treated 87 cases of lumbar spinal stenosis with percutaneous endoscopic technology. The results showed that the VAS, ODI, and claudication distance were significantly improved compared with baseline. A significant difference in efficacy was observed between the 71–81 year age group and the other age groups. The actual spinal canal area was positively correlated with age. The soft tissue invasion ratio of the vertebral canal and invasion ratio of the nerve root canal were negatively correlated with age. There was a correlation between the soft tissue

### Table 3 Correlation analysis between sex and CT/MRI parameters

|                        | r     | P    |
|------------------------|-------|------|
| Bony vertebral canal area by CT measurement | –0.008 | 0.939 |
| Cross-sectional area of the dural sac by MRI measurement | –0.162 | 0.133 |
| Nerve root canal bony area by CT measurement | –0.014 | 0.895 |
| Yellow ligament thickness by MRI measurement | –0.010 | 0.927 |
| Real spinal canal area by CT measurement | 0.223 | 0.038 |
| Nerve root canal real area by CT measurement | 0.125 | 0.249 |
| Vertebral canal sagittal diameter by CT measurement | –0.041 | 0.707 |
| Distance between the articular joints by CT measurement | 0.007 | 0.948 |
| Soft tissue invasion ratio of the vertebral canal | –0.228 | 0.033 |
| Invasion ratio of the nerve root canal | –0.360 | 0.001 |

### Table 4 Correlation analysis between age and CT/MRI parameters

|                        | r     | P    |
|------------------------|-------|------|
| Bony vertebral canal area by CT measurement | –0.012 | 0.349 |
| Cross-sectional area of the dural sac by MRI measurement | –0.607 | 0.538 |
| Nerve root canal bony area by CT measurement | –0.025 | 0.820 |
| Yellow ligament thickness by MRI measurement | 0.015 | 0.893 |
| Real spinal canal area by CT measurement | 0.188 | 0.081 |
| Nerve root canal real area by CT measurement | 0.080 | 0.463 |
| Vertebral canal sagittal diameter by CT measurement | –0.170 | 0.115 |
| Distance between the articular joints by CT measurement | 0.047 | 0.667 |
| Soft tissue invasion ratio of the vertebral canal | –0.211 | 0.049 |
| Invasion ratio of the nerve root canal | –0.215 | 0.045 |
invasion ratio of the vertebral canal and the invasion ratio of the nerve root canal with sex. Correlation analysis of the imaging parameters and effect revealed a good correlation between the clinical efficacy and vertebral canal sagittal diameter, distance between the articular joints, soft tissue invasion ratio of the vertebral canal, and invasion ratio of the nerve root canal. These measurements were positively correlated with VAS, ODI, and claudication distance.

**Conclusion**

Based on the above results, the following conclusions can be drawn. The treatment of lumbar spinal stenosis by endoscopic transforaminal decompression can achieve good clinical results, but the operation could be less effective in patients > 71 years of age. Whether age > 71 is a contraindication of treatment of lumbar spinal stenosis with percutaneous endoscopic technology remains to be confirmed. There was a positive correlation between the vertebral canal sagittal diameter, distance between the articular joints, soft tissue invasion ratio of the vertebral canal, and invasion ratio of the nerve root canal with the clinical effect. This is in line with the characteristics of the intervertebral foramen mirror because of the greater influence factors, such as age.

In conclusion, although percutaneous endoscopic decompression surgery can have good clinical efficacy and unique minimally invasive advantages in the treatment of degeneration of spinal canal stenosis [15–18], the surgical indications are dependent on a good preoperative evaluation because the intervertebral foramen is still a greater challenge than lumbar intervertebral disc herniation. It is also important to focus on the complications of surgery such as dural sac and nerve injury [19, 20], as well as radiation exposure [21]. Due to the limited amount of clinical data collection and limited follow-up time, the results of this study have a clear guiding significance for the clinical application of intervertebral endoscopy in the treatment of lumbar spinal stenosis, which requires further study and exploration.

| Table 5 | Correlation analysis between CT/MRI parameters and therapeutic effects |
|---------|-------------------------------------------------------------------------|
|         | VAS | ODI | Walking distance |
|         | r   | P   | r   | P   | r   | P   |
| Bony vertebral canal area by CT measurement | 0.166 | 0.124 | 0.167 | 0.122 | 0.136 | 0.210 |
| Cross-sectional area of the dural sac by MRI measurement | 0.112 | 0.304 | 0.024 | 0.826 | 0.195 | 0.071 |
| Nerve root canal bony area by CT measurement | −0.119 | 0.272 | 0.035 | 0.750 | 0.001 | 0.992 |
| Yellow ligament thickness by MRI measurement | 0.162 | 0.133 | 0.012 | 0.909 | 0.122 | 0.260 |
| Real spinal canal area by CT measurement | −0.041 | 0.705 | −0.102 | 0.348 | 0.044 | 0.687 |
| Nerve root canal real area by CT measurement | −0.179 | 0.098 | −0.103 | 0.344 | 0.008 | 0.943 |
| Vertebral canal sagittal diameter by CT measurement | 0.139 | 0.198 | 0.050 | 0.643 | 0.055 | 0.613 |
| Distance between the articular joints by CT measurement | 0.366 | < 0.001 | 0.222 | 0.039 | 0.377 | < 0.001 |
| Soft tissue invasion ratio of the vertebral canal | 0.228 | 0.021 | 0.213 | 0.047 | 0.261 | 0.024 |
| Invasion ratio of the nerve root canal | 0.262 | 0.014 | 0.290 | 0.006 | 0.350 | 0.001 |

| Table 6 | Linear regression of different indicators (multiple linear regression) |
|---------|---------------------------------------------------------------|
| Parameters | VAS (6 months) | ODI (6 months) | Walking distance (6 months) |
|          | Non-standardized β | Standardized β | P     | Non-standardized β | Standardized β | P     | Non-standardized β | Standardized β | P     |
| Soft tissue invasion ratio of the vertebral canal | 71.05 | 0.542 | < 0.001 | 31.592 | 0.375 | < 0.001 | 2296.6 | 0.435 | < 0.001 |
| Invasion ratio of the nerve root canal | 28.06 | 0.347 | 0.015 | 6.21 | 0.025 | 0.04 | |
| Real spinal canal area by CT measurement | 0.3 | 0.023 | 0.011 | |
| Nerve root canal real area by CT measurement | −0.08 | 0.01 | 0.034 | |
| Walking distance | 635.4 | 0.376 | 0.028 | | | | | | | |
Supplementary information

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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