Patient Reported Outcome Analysis after Posterior Decompression and Posterolateral Fusion Surgery in Lumbar Degenerative Spondylolisthesis

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Abstract

Introduction: Lumbar degenerative spondylolisthesis is an acquired slippage of a vertebra due to degenerative arthritis of the facet joints without any defect in the vertebral ring. Degenerative spondylolisthesis is commonly treated with posterior decompression and posterolateral fusion. This study aims to evaluate the clinical outcome of posterior decompression and instrumented posterolateral fusion surgery by analysing patient reported outcome measures.

Materials and Methods: A retrospective study in which patients who had posterior decompression and instrumented posterolateral fusion for lumbar degenerative spondylolisthesis between September 2017 and August 2019 at Nepal Mediciti Hospital were included in the study. Other types of spondylolisthesis managed with similar technique and patients who had follow up period of less than three months were excluded. Functional outcome was assessed by comparing pre and post-operative patient reported outcome measures: VAS leg pain, VAS back pain and Oswestry Disability Index (ODI). The paired t-test was used for statistical analysis.

Results: Of the 16 patients, 81.25% were female. Mean age and mean follow up period were 58.81± 10.47 years and 13.56±7.15 months respectively. Seventy five percent had grade I spondylolisthesis. Most common level of spondylolisthesis was L4/5 (62.5%). Fourteen patients reported improvement in their symptoms after surgery. The changes in functional outcome scores between baseline and at follow up evaluation were as follows: 32.70 ± 17.44 points for ODI, 3 ±1.89 for VAS leg pain, and 5.37 ± 2.36 for VAS back pain (P<0.001). Superficial wound infection was the most common complication observed in 18.3%.

Conclusion: In our case series, 87.5% had improvement in their symptoms after surgery. The change in mean score of patients’ reported outcome measures before and after surgery was statistically significant. We recommend a prospective comparative study between decompression alone and the fusion technique for the evaluation of long term functional outcome.

Key words: Degenerative Spondylolisthesis, Lumbar, Instrumented Posterolateral Fusion, Outcome

Introduction

Degenerative spine disease (DSD) is one of most common spine pathology affecting up to 80% of the population at some point in their life. The annual

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Prevalence of DSD is 15% to 40% leading to disability in 1% to 2% of the population. Spinal stenosis and spondylolisthesis, the sequelae of degeneration, together affect up to 20% of the U.S. population.

Newman and Stone coined the term degenerative spondylolisthesis (DS) in 1955, who defined it as an acquired slippage of the vertebra secondary to degenerative arthritis of facet joints without defect in pars interarticularis. Karkaldy Willis described the degenerative cascade in 1970s. The cascade begins with disc degeneration, progresses to destabilization and finally leads to re-stabilization. The cause of pain in DS has been described as multi-factorial: purely mechanical pain due to degenerative changes or neurogenic claudication from spinal stenosis or radicular pain due to nerve root impingement in the lateral recess or neural foramen.

In this spectrum of cascade, surgery is indicated for instability, progressive neurological deficit or those not responding to medical management.

Although various surgical techniques have been used for the management of DS, the superiority of one over other was not well evident in the literature. In general, outcomes were commonly assessed based on surgeons’ perspective relying on technical success like adequate decompression and/or fusion. Interestingly, surgeons’ perspectives and results of imaging exams frequently do not found to be correlated with patient satisfaction. In this context, considering the importance of patient’s opinion of results and their satisfaction, we designed a study to evaluate the outcome of posterior decompression and instrumented posterolateral fusion (PLF) surgery in patients with lumbar DS using patient reported outcome instruments—visual analogue scale (VAS) leg pain, VAS back pain and Oswestry Disability Index (ODI).

Materials and Methods

Study Design and Patient Population

We designed a retrospective study. Ethical approval was taken from Institutional Review Committee of the hospital. A consecutive cohort of patients who had posterior decompression and instrumented posterolateral fusion (PLF) surgery for lumbar degenerative spondylolisthesis between September 2017 and August 2019 at Nepal Mediciti Hospital was identified from medical record files. Patients whose data were incomplete, who had follow up period of less than three months and all other types of spondylolisthesis were excluded. Demographic, clinical, radiographic reports and operative records were reviewed. Patients were evaluated with respect to their age, sex, duration of symptoms, clinical presentation, level and grade of spondylolisthesis. Grading of spondylolisthesis was classified from grade I to V according to Meyerding classification system. Any intra-operative and/or post-operative complication was documented.

Pre-operative management and surgical indication

All patients were managed with conservative treatment with adequate analgesics and physical therapy for at least 12 weeks. Indication for grade I degenerative spondylolisthesis was failure of conservative management and unstable spondylosis (translation in spondylolisthesis in standing lateral flexion/extension x ray of lumbosacral spine) while the indication for surgery in grade II spondylolisthesis was failure of conservative treatment with progressive neurological deficit.

Surgical technique

Surgical procedure was performed through an open midline longitudinal approach in prone position. Pedicle screws were placed based on anatomic landmarks and fluoroscopic image assistance. The neural element decompression was performed by a midline laminectomy, medial facetectomy and foraminotomy. Posterolateral fusion was facilitated by inter-transverse grafting along the decorticated surface of the transverse process using autologous local bone graft collected during laminectomy. Post-operative antero- posterior and lateral X ray of lumbosacral region was done to confirm the position of implants and alignment of bony elements.

Study Measures/Statistics

Outcome data included visual analogue pain scale (VAS) and Oswestry Disability Index (ODI). Patients were consented and asked to complete a survey before surgery and at the time of follow up at least three months after surgery in out-patient department. Outcome was assessed by comparing pre and post-operative patient-reported outcome measures: visual analogue scale (VAS) leg pain, visual analogue scale (VAS) back pain and Oswestry disability index (ODI). The paired t-test was used for statistical analysis. SPSS 17 (IBM, Chicago, IL, USA) was used for data analysis. P value <0.05 was considered significant.

Results

Of 19 patients with degenerative lumbar spondylolisthesis were operated in 24 months, 16 were included in the study. Patient demographics and clinical characteristics are summarized in Table 1. Majority of the patients were female (81.25%), with female to male ratio of 4.3:1. Mean age was 58.81 years (SD 10.47, range 45-80). The mean and the
minimum duration of follow up were 13.5 and 5 months respectively. 75% had grade I spondylolisthesis while grade II spondylolisthesis was seen in 25%. Most of the deformity was seen at L4/5 Level followed by L5/S1 level (62.5% vs 38.5%).

87.5% improved after surgery. The changes in functional outcome scores between baseline and at postoperative follow up evaluation were as follows: 32.70 ± 17.44 points for ODI, 3 ±1.89 for VAS leg pain, and 5.37 ± 2.36 for VAS back pain. Mean scores of VAS leg pain, VAS back pain and ODI before surgery and during follow up were presented in Table 2.

Most common perioperative complication in our series was wound dehiscence seen in 18.75% followed by intra-operative cerebrospinal fluid (CSF) leak (6.5%). All the patients who had wound dehiscence in our series (n=3) were elderly diabetic (age > 65 years). One of them was suffering from Cushing disease who underwent two level fixation and fusion because of risk of hypercortisolemia-induced osteoporosis. (See Figure 1 and 2)

![Figure 1](image1.png)

*Figure 1:* (A) MRI image of lumbosacral Spine of 45-year female with Cushing disease showing grade II degenerative spondylolisthesis at L5/S1 level, (B) Post operative X ray images showing reduction with instrumented posterolateral fixation and fusion.

![Figure 2](image2.png)

*Figure 2:* (A) MRI Lumbosacral Spine of 80-year female showing grade I degenerative spondylolisthesis at L4/5 level, (B) Post-operative X ray images showing reduction with instrumented posterolateral fixation and fusion. We included L5/S1 in this case due to old age and osteopenic bone to avoid junctional failure.
Figure 3: Evolution of Treatment paradigm in Degenerative Spondylolisthesis

Table 1: Patient Baseline Demographic and Clinical Characteristics, N=16

| Variables                        | Pre-operative | Post-operative | Mean difference (95% CI) | P Value |
|---------------------------------|---------------|----------------|--------------------------|---------|
| VAS leg pain (mean ± SD)        | 6.81±1.56     | 3.81±1.22      | 3.0 ±1.89 (1.98-4.01)    | <0.001  |
| VAS back pain (mean ± SD)       | 8±1.03        | 2.63±1.85      | 5.37±2.36 (4.11-6.63)    | <0.001  |
| ODI (mean ± SD)                 | 56.66±10.33   | 23.96±12.40    | 32.70±17.44 (23.41-42)   | <0.001  |

Table 2: Comparison of Mean Change in Outcomes before and after PLF (Posterolateral fusion)

Discussion

Previous studies have shown that degenerative spondylolisthesis is commonly seen in elderly female (above 50 years) with a female-to-male ratio of 6:1, with L4/5 being the most common level of deformity. This was consistent with our findings.

Meyerding classification is widely used to classify DS. Low grade spondylolisthesis is the commonest deformity associated with degeneration. As consistent with previous studies, all of our patients had low grade spondylolisthesis (grade I and II) with grade I being the most common entity (75%).

In general, a minimum of 2-year follow-up is suggested for patient-reported outcomes (PROs) to

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adequately assess the therapeutic effect of surgery in spine and musculoskeletal clinical research. Ayling et al investigated on the appropriate follow up time for clinical outcome research in spine surgery and found that the time to plateaued recovery after surgery for VAS back and leg pain was 3 months; and 12 months for disability indices (ODI) for patients with degenerative spondylolisthesis who were treated with fusion. In our study the minimum and the mean duration of follow up were 5 and 13.5 months respectively.

In our study, all patients were managed with PLF. Fourteen out of sixteen patients improved after surgery. There was significant change in the means of VAS leg pain, back pain and ODI scores from 6.8, 8, and 56.66 before surgery to 3.8, 2.6 and 23.96 respectively after surgery (p<0.001). Omidi-Kashani F et al published a similar study in which pre-operative pain and disability (VAS and ODI) improved from 8.8 and 71.6 to postoperative 2.1 and 28.7 respectively after PLF surgery lumbar DS. Urquhart et al did a retrospective study and found that the change in mean of leg pain, back pain and ODI scores after PLF for lumbar DS were -5.9, -4.1 and -26 respectively, which was similar to our findings in which the change in the means were -3, -5.37 and -32.7 respectively. Similarly, Gottschalk et al found a significant change in visual analogue pain scale (MCID) of 3 months; and 12 months for disability indices (ODI). The minimum difference in score in our study were 15.26 for ODI, 1.11 for back pain, and 3.01 for leg pain which correspond to the MCID reported in literature.

In our case series, 87.5% had improvement in their symptoms after surgery. The change in mean score of patients’ reported outcome measures before and after surgery was statistically significant (P<0.001). Based on patient reported outcomes, PLF seems to be a viable treatment option in degenerative spondylolisthesis after laminectomy alone in patients with stable grade I DS. Hence fusion or no fusion and the technique of fusion are the topics of ongoing discussion and the treating surgeon has to tailored the approach based on clinical scenario, available evidences and the patient’s decision. In our case series, we performed fusion surgery in grade I DS because of translation seen in standing dynamic X ray of lumbosacral spine while in cases of grade II DS fusion was recommended due to high risk of progression of spondylolisthesis with laminectomy alone. The evolution and controversy in the management of lumbar degenerative spondylolisthesis was summarized in figure 3.

Our perioperative complications were higher than that of the findings (wound dehiscence- 6.9% and CSF leak-4.6%) reported by Urquhart et al. Post-operative wound infection and dural tear were seen in 10% and 11% respectively in SSSS trial.

Limitation

This study was limited by small sample size and its retrospective nature. We were not able to assess the evidence of radiological fusion post-operatively. Though smaller in size, this study is trying to incorporate patient’s perspective regarding outcome of surgical intervention after spinal surgery in Nepal. Patient’s satisfaction and cost effectiveness are important considerations while selecting the surgical technique especially in the context of developing country like Nepal where affordability is a major concern.

Conclusion

In our case series, 87.5% had improvement in their symptoms after surgery. The change in mean score of patients’ reported outcome measures before and after surgery was statistically significant (P<0.001). Based on patient reported outcomes, PLF seems to be a viable treatment option in degenerative spondylolisthesis. We recommend a prospective comparative study between decompression alone and the fusion technique to assess the long-term functional outcome.

Conflict of Interest: None

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