Keywords: Discectomy; Lumbar disc; MRI imaging; Antibiotics

Introduction

Backache is one of the most common clinical symptoms encountered in medical practice. Low back pain is a symptom, not a disease. The pathologic basis for the pain may be something within the spine or a lesion outside of the spine [1] up to 80% of patients with acute low back pain, a precise anatomic cause cannot be localized. Typically, arrays of non-specific terms are used like lumbago, sprain, strain, mechanical low back pain and lumbar syndrome.

Prolapsed intervertebral disc is an important cause of spondylogenic backache. Although back pain is common from the second decade of life, intervertebral disc disease and disc herniation are most prominent in otherwise healthy people in third and fourth decades of life. Ninety five percent of lumbar disc herniation occurs at either L4-L5 or L5-S1 level [2].

The posterior longitudinal ligament affords only weak reinforcement especially at L4-L5 and L5-S1 level where it is a midline narrow unimportant structure attached to annulus [3].

The L4-L5 and L5-S1 articulations have the greatest motion in the lumbar spine. Greater motion causes an increased potential for instability, degeneration and breakdown and therefore the incidence of herniated discs is greater at L4-L5 and L5-S1 level than at any other lumbar disc space [4].

A Visual Analog Scale (VAS) [5] is a measurement instrument that tries to measure a characteristic or attitude that is believed to range across a continuum of values and cannot easily be directly measured.

Abstract

Background: This study was to find out that whether the lumbar disc prolapse at different level influence the functional outcome of patients after discectomy. Present study is of 50 adult cases admitted at D. Y Patil Medical College. Common age group involved was between 40-60 years.

Aims: To assess whether there is any difference in functional outcome of patients with disc prolapse at different levels in the lumbar spine after performing discectomy.

Objectives: To compare pre-operative and post-operative:
- Leg pain and back pain by Visual Analog Scale Score
- Functional outcome by Modified Oswestry Disability index score of the selected patients.

Materials and methods: This study was a comparative study, conducted for a period of about two years, July 2014 to September 2016 in Dr. D. Y. Patil Medical College, Hospital & Research Centre, only patients who were scrutinized for exclusion criteria and also abiding to inclusion criteria were included. Period required for data collection: 2 years. Period required for data analysis and reporting: - 6 months. We prospectively followed 50 consecutive patients with unilateral lumbar herniation either at L2-L3, L3-L4, L4-L5 or L5-S1 levels requiring surgery. The procedure performed was Micro lumbar discectomy in all patients.

Results: A total of 50 patients were included in our study of which 4 (8%) patients had prolapsed intervertebral disc at L2-L3 level and 10 (20%) patients had disc prolapsed at L3-L4 level. These 14 patients were included in upper lumbar level disc herniation group referred henceforth as Group 1 (28%). 22 (44%) patients had disc prolapsed at L4-L5 level and 14 (28%) patients had disc prolapsed at L5-S1 level, these 36 patients were included in lower lumbar level disc herniation group who are referred henceforth as Group 2 (72%).

On comparing the results after discectomy of prolapsed intervertebral disc at different levels in the lumbar spine we found no significant difference in the end result and functional outcome of the patients.

Conclusion: The aim of this study was to find out that whether the lumbar disc prolapse at different levels influence the functional outcome of patients. According to the observations of this study and after reviewing various similar studies done in the past we conclude that after discectomy, level of disc prolapse per se has no significant bearing on functional outcome of the patients.
For example, the amount of pain that a patient feels ranges across a continuum from none to an extreme amount of pain.

The Modified Oswestry Disability Index (ODI) \cite{6,7} is an extremely important tool that researchers used to measure patient's functional disability due to low back pain. The test is considered as gold standard of low back functional outcome tools. The ODI is a valid and vigorous measure and has been a worthwhile outcome measure. It consists of questionnaire which has been designed to give us information as to know how the back pain has affected patient's ability to manage in everyday life. It categories the patient as having minimal, moderate or severe disability due to back pain.

**Materials and Methods**

This study is a comparative study was conducted for a period of about two years, July 2014 to September 2016 in Dr. D. Y. Patil Medical College, Hospital & Research Centre. Only patients who were scrutinized for exclusion criteria and also abiding to inclusion criteria were included. Period required for data collection: 2 years. Period required for data analysis and reporting: 6 months. A study done by Murthy H, Reddy TVS showed a significant reduction in mean preop vas score from 7.47 to 1.8016 with power of study 80% so We prospectively followed 50 consecutive patients with unilateral lumbar herniation at either L2-L3, L3-L4, L4-L5 or L5-S1 levels requiring surgery. All patients included in this study were operated by the same orthopaedic surgeon in the same operation theatre as the first case of the day. The procedure performed was Microlumbar discectomy in all patients \cite{8}. Minimum follow up in this study was 2 months and maximum follow up was 20 months. Consecutive patients of either sex, in the age group 20yrs - 55yrs, who fulfilled the understated criteria, were operated. A total of fifty patients were selected with following inclusion criteria \cite{3}.

**Inclusion criteria**

- Patients over eighteen years old who had radicular pain for atleast four weeks with positive nerve root tension sign & who had no relief after non-operative treatment like bed rest, analgesics, and traction for 4 weeks.
- Confirmatory cross-sectional MRI imaging study demonstrating intervertebral disc herniation at a level and side corresponding to their symptoms.
- Patients lying in ODI scoring group III were only included
- All patients who will have follow up of at least 3 months will be part of this study

Standard open fenestration and discectomy: Preoperative preparations: Patient was kept nil orally since the night prior to the day of operation. Entire back was prepared by shaving the part and thorough soap and water wash was given. Preoperative antibiotics were administered. Patient was induced by general anesthesia. The patient was placed prone in the knee-chest position. The abdomen was kept free, so as to keep the respiration free and prevent engorgement of the epidural veins and thus reduce bleeding (Figure 1).

**Approach**

A mid-line vertical incision over the affected interspace of 8-10 cm is made after the back has been thoroughly painted and draped. The incision is deepened to the subcutaneous tissue and deep fascia. The lumbodorsal fascia is incised and the supraspinous ligament is incised over the affected disc space. By subperiosteal dissection, strip the paraspinal muscles from the spines and laminae of the vertebrae on each side and self-retaining retractors are applied. Using microscope partial laminectomy and discectomy was done using routine surgical technics (Figure 2).

**After care**

Patient was allowed to turn in bed. Pain was controlled with injectable and oral NSAIDS. Postoperative antibiotic were administered. 24 hours post operatively patient was again examined for the severity of back pain and leg pain VAS score for this were noted (Figure 3).
Neurological function was closely monitored after surgery. Neurological function was monitored closely. For urinary retention patients were given antispasmodics and encouraged to pass urine. Catheterization was done if supportive measures failed. Sutures were removed after 10-12 days (Figure 4).

Follow up evaluation
- The leg and back pain analysis by VAS score was done 24 hours after surgery, then every week for 3 weeks, then at 1st month, 2nd months and finally at 3rd months on follow-up.
- The ODI was evaluated at 1st month, 2nd month and then at 3rd months after the surgical procedure.
- These findings were compared between the groups for difference if any by statistical analysis.

Results
In our study patient data was recorded pre-operatively, 24 hrs after surgery then on follow-up for every week for initial one month and then monthly for three months. Patients was called and evaluated on objective and subjective criteria, objectively, we use straight leg raising test [8], femoral stretch test [6] and Lasegue test. Subjectively we use visual analogue test, The Modified Oswestry Disability Index (ODI) [6,7] (Table 1).

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Group I (n=14)</th>
<th>Group II (n=36)</th>
<th>p-value</th>
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<td>Mean</td>
<td>SD</td>
<td>Mean</td>
<td>SD</td>
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<tr>
<td>Age (yrs)</td>
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<td>42.28</td>
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<td>Gender (M:F)</td>
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<td>27:9</td>
<td>0.847</td>
</tr>
</tbody>
</table>

Table 1: Level wise distribution of patients.

A total of 50 patients were included in our study of which 4 (8%) patients had prolapsed intervertebral disc at L2-L3 level and 10 (20%) patients had disc prolapsed at L3-L4 level. These 14 patients were included in upper lumbar level disc herniation group referred hereafter as Group 1 (28%). 22 (44%) patients had disc prolapsed at L4-L5 level and 14 (28%) patients had disc prolapsed at L5-S1 level. These 36 patients were included in lower lumbar level disc herniation group who are referred hereafter as Group 2 (72%) (Table 2).

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<td>9:5</td>
<td>27:9</td>
<td>0.847</td>
</tr>
</tbody>
</table>

Table 2: Comparison of age and gender in group I and group II.

Out of all the patients in the study a total of 36 (72%) were males and 14 (28%) were females. In Group 1 i.e., upper lumbar level patients, 9 (64.29%) patients were male and 5 (35.71%) patients were female. In group 2 i.e., lower lumbar level patients, 27 (75%) were males and 9 (25%) were females. By using 2 samples proportion tests, p-value>0.05, therefore there is no significant difference between the proportion of gender in group 1 and group 2 (Table 3).

<table>
<thead>
<tr>
<th>Gender</th>
<th>Number of patients</th>
<th>Percentage (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male</td>
<td>36</td>
<td>72</td>
</tr>
<tr>
<td>Female</td>
<td>14</td>
<td>28</td>
</tr>
<tr>
<td>Total</td>
<td>50</td>
<td>100.00</td>
</tr>
</tbody>
</table>

Table 3: Gender wise distribution of patients.

The mean age of patients in the group 1 was 55.21 years (SD± 6.10) with a minimum age of 48 and a maximum age of 70 while in the group 2, mean age being 42.28 years (SD± 10.80) with minimum age
of 20 and a maximum of 59. By using 2 independent sample t-test, p-value<0.05, therefore there is significant difference between group 1 and group 2 patients with respect to age (years) (Table 4).

The maximum number of patients i.e., 8 in group 1 are in 51-60 yrs age group while in group 2, 18 patients are in 41-50 yrs age group. Thus, we conclude that upper lumbar level disc herniation is more common in elderly age group while lower lumbar level disc prolapse is more common in middle age group (Table 5).

### Table 5: Age wise distribution in Group 1 and 2.

Visual analog scale was used to quantify leg pain and back pain. The mean VAS score for leg pain in the group 1 in the pre-operative period was 9.40 (SD=± 0.87) with 1 (7.14%) patient having minimum score of 7 and 4 (28.57%) patients had maximum score of 10. In group 2 the mean VAS score for leg pain was 2.79 (SD=± 0.69) with 5 (35.71%) patients having minimum value of 7 and 23 (63.88%) patients had maximum value of 10. There was no significant difference (p-value=0.121) in preoperative VAS scores for leg pain in group 1 and group 2 (Table 6).

### Table 6: Comparison of age (years) in group 1 and 2.

At 24 hrs post-operatively the mean leg pain in the group 1 was reduced to 2.79 (SD=± 0.69) with 5 (35.71%) patients having minimum of 2 and a maximum score of 4 was noted in 2 patients (14.28%) whereas in the group 2 the mean score was reduced to 1.14 (SD=± 0.74) with a minimum value of 0 and a maximum score of 3 was recorded in 1 (7.14%) patient. The difference in VAS scores for pain at 24 hrs post-operatively between group 1 and 2 was not significant (p-value=0.074).

On follow at one week the mean leg pain was 0.71 (SD=± 0.61) with 5 (35.71%) patients had no pain (VAS=0) and single patient (7.14%) had maximum score 2 in group 1 whereas in the group 2 mean score for leg pain was 2.89 (SD=± 0.89) with no pain in 15 (41.66%) patients (VAS=0) and a maximum score of 4 was found in single patient (2.77%). The difference in VAS scores for pain at one week follow up between group 1 and 2 was not significant (p-value=0.682).

However, at two weeks follow up the mean leg pain was 0.43 (SD=±0.51) in the group 1 while in the group 2 mean leg pain was 0.11 (SD=±0.40). At three weeks follow up only two patients in group 1 had leg pain while in group 2 no patient reported leg pain. On further follow up all patients in both groups reported no leg pain.

By using 2 independent sample t-test p-value>0.05, pre-operatively, 24 hr post-operative, at 1st week to 3rd month. Therefore, we conclude that there is no significant difference between the mean VAS score for leg pain pre-operatively, 24 hrs post-operatively, at 1st week, 2nd week, 3rd weeks, 1st month, 2nd month and 3rd month follow-up in group 1 and group 2 (Table 7).

### Table 7: Comparison of VAS score for leg pain in group 1 and 2.

The mean VAS score for back pain in group 1 in the pre-operative period was 4.52 (SD=± 1.57) with 1 (2.77%) patient having minimum score of 2. However, at two weeks follow up the mean leg pain was 0.43 (SD=±0.51) in the group 1 while in the group 2 mean leg pain was 0.11 (SD=±0.40). At three weeks follow-up only two patients in group 1 had leg pain while in group 2 no patient reported leg pain. On further follow up all patients in both groups reported no leg pain.

By using 2 independent sample t-test p-value>0.05, pre-operatively, 24 hr post-operative, at 1st week to 3rd month. Therefore, we conclude that there is no significant difference between the mean VAS score for leg pain pre-operatively, 24 hrs post-operatively, at 1st week, 2nd week, 3rd weeks, 1st month, 2nd month and 3rd month follow-up in group 1 and group 2 (Table 7).

### Table 8: Comparison of VAS scores for back pain in group 1 and 2.

The mean VAS score for back pain in group 1 in the pre-operative period was 6.57 (SD ± 1.22) with 3 (21.42%) patient having a minimum score of 1 and a maximum of 9 which was noted in 1 (7.14%) patient. In group 2 the mean pre-operative VAS score for back pain was 2.29 (SD=± 0.99) with 0 (0.0%) patient having a minimum score of 0 and a maximum of 2 (14.28%) with maximum of 8. There is significant difference (p-value=0.019) in preoperative VAS scores for back pain in group 1 & group 2.
back pain 24 hours after surgery was 1.58 (SD=± 1.59) with minimum score of 0 noted in 12 (33.33%) patients and a maximum of 5 in 2 (5.55%). There Was significant difference (p-value=9.008) in post-operative 24 hours VAS scores for back pain in group 1 and group 2 (Table 8).

On follow at one week the mean leg pain was 0.86 (SD=± 0.95) with a minimum score of 0 noted in 7 (50%) patients and a maximum of 5 in 2 (14.28%) patients and a maximum score of 4 in 1 (2.77%) patient. The difference in VAS scores for back pain at one week follow up between group 1 and group 2 was NOT significant (p-value=0.554).

However, at two weeks follow up the mean back pain was 0.36 (SD=0.63) in the group 1 while in the group 2 was 0.17 (SD=0.74). On further follow up patients in group 1 had some back pain while patients in group 2 had no back pain except for one patient who had complication of wound infection.

By using 2 independent sample t-test, p-value <0.05 pre-operatively and 24 hrs post-operatively but p-value>0.05, at follow up from 1st week to 3rd month post operatively. Therefore, we can conclude that there is no significant difference between the mean VAS score for back pain at 1st week, 2nd week, 3rd week, 1st month, 2nd month and 3rd month in group 1 and group 2 though there is significant difference preoperatively and initially post-operatively at 24 hrs. Back pain was more significant in upper lumbar disc prolapses (Table 9).

### Table 9: Comparison of Oswestry score in group 1 and 2.

<table>
<thead>
<tr>
<th>Oswestry score</th>
<th>Group I (n=14)</th>
<th>Group II (n=36)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean</td>
<td>SD</td>
<td>Mean</td>
<td>SD</td>
</tr>
<tr>
<td>Pre-operative</td>
<td>52.29</td>
<td>4.76</td>
<td>50.28</td>
</tr>
<tr>
<td>Post op 1st month</td>
<td>10.43</td>
<td>1.95</td>
<td>20.94</td>
</tr>
<tr>
<td>Post op 2nd month</td>
<td>6.71</td>
<td>2.01</td>
<td>12.11</td>
</tr>
<tr>
<td>Post op 3rd month</td>
<td>5.43</td>
<td>1.45</td>
<td>6.28</td>
</tr>
</tbody>
</table>

We used modified Oswestry Disability questionnaire and calculated the Oswestry Disability Index pre-operatively and at follow-ups at 1st month, 2nd months and 3rd months post-operatively to evaluate the functional outcome of patients. The pre-operative mean ODI in group 1 was 52.29 (SD=4.76) while that in group 2 was 50.28 (SD=6.01). However, there was no significant difference (p-value=0.225) in the pre-operative ODI in the two groups.

The mean ODI in group 1 reduced to 10.43 (SD=1.95) at 1 month follow up while that in group 2, the mean ODI reduced to 20.94 (SD=4.76) at 2nd week, 3rd week, 1st month, 2nd month and 3rd month respectively. However, there was no significant difference between the two groups (p-value=0.001). The difference in the ODI scores between the two groups was significant (p-value=0.001). At 3rd month of follow up the mean ODI in group 1 reduced to 5.43 (SD=1.45) while that in group 2 reduced further to 6.28 (SD=1.53). This difference in the ODI scores between the two groups was NOT significant at third month of follow up (p-value=0.079).

By using 2 independent sample t-test p-value>0.05 therefore we conclude that there is no significant difference between the mean Oswestry score at pre-operatively and at 3rd month follow-up in group 1 and group 2 but initially at 1st month and 2nd month follow up there is significant difference in Oswestry scoring in two groups.

Patients with upper or higher lumbar level lesions improved significantly initially after discectomy with better functional outcome but on long term follow up functional outcomes in both groups after discectomy were same.

All patients in both groups, as a protocol, were discharged on the 5th postoperative day. Only one patient in group 2 had complication of wound infection. No other complications were noted. All patients by the end of follow up (3rd month) had returned to their original job.

### Discussion

We prospectively followed 50 consecutive patients with unilateral lumbar herniation at L2-L3, L3-L4, L4-L5 or L5-S1 levels after discectomy.

### Level of disc prolapse

A total of 50 patients which were included in our study of which only 4 (8%) patients had disc prolapse at L2-L3 level while 10 (20%) patients had disc prolapse at L3-L4 level who are referred as Group 1. L2-L3 and L3-L4 levels constitutes so called upper lumbar disc herniation group. 22 (44%) patients had prolapsed intervertebral disc at L4-L5 level and 14 (28%) at L5-S1 level, who are referred as Group 2 [9]. L4-L5 and L5-S1 levels constitutes so called lower lumbar disc herniation group. Thus, most common level of disc prolapse in our study is L4-L5 level followed by L5-S1 level which together forms 72% of our patients. According to literature, the majority of lumbar herniation occurs at the L4-L5 and L5-S1 intervertebral disc levels, affect the L5 and S1 roots, and result in sciatica. Upper level herniation (levels L2-L3 or L3-L4) are less common, may affect the L2, L3, and L4 nerve roots and cause a femoral radiculopathy [10-13].

### Sex distribution

Out of all the patients in the study a total of 36 (72%) were males and 14 (28%) were females. In Group 1 that is upper lumbar level patients, 9 (64.29%) patients were male and 5 (35.71%) patients were female. In group 2 that is lower lumbar level patients, 27 (75%) were males and 9 (25%) were females. By using 2 samples proportion test p-value>0.05 therefore there was no significant difference between the proportion of gender in group 1 and group 2 in our study. But disc problem was more preponderant in males than females. Similarly, in the study done by Lurie and et al., the majority of the study population (57%) was male [14]. Also in study done by Saberi et al., male to female ratio was 1.08 and 1.14 in the upper and lower lumbar disc herniation, respectively [15].

### Age significance

The mean age of patients in the group 1 was 55.21 years (SD± 6.10) with a minimum age of 48 and a maximum age of 70 while in the group 2, mean age being 42.28 years (SD± 10.80) with minimum age of 20 and a maximum of 59. By using 2 independent sample t-test p-values<0.05, therefore there was significant difference between group 1 and group 2 patients with respect to age (years). The maximum number of patients i.e., 8 in group 1 were in 51-60 years of age while in group 2, 18 patients were in 41-50 years age group. Thus, we conclude that upper lumbar level disc herniation was more common in elderly age group while lower lumbar level disc prolapse was more common in
middle age group. In the study done by hooshang Saberi et al., The mean age of patients with upper lumbar disc herniation and lower lumbar disc herniation were 45.7 years (23-70) and 41.2 years (20-63), respectively.43. Similarly, in SPORT trial by Lurie et al. case study the level of herniation varied directly with age, as patients with upper level herniation were significantly older, the L4-L5 group was of an intermediate age, and the L5-S1 group was the youngest [14].

Pain analyses

The severity of the leg pain and back pain was noted by the VAS score. The pre-operative VAS scores were more for leg pain than back pain in both groups. Patients in both groups had more leg pain than back pain. Patients in group 1 with upper lumbar level disc prolapse had more back pain as compared to patients in group 2 with lower lumbar level disc prolapse.

Leg pain:

1. Pre-operative leg pain in group 1 (mean VAS=9.4) was not significantly different (p-value=0.12) from the leg pain in group 2 (mean VAS=9.44). There was an initial rapid decrease in the leg pain scores in both groups from 9.4 in the pre-operative period to 2.79 and 2.89 respectively 24 hrs after operation. This rapid decrease in leg pain scores is similar in both groups (p-value=0.74).

2. On follow at one week the mean leg pain was 0.71 (SD=± 0.61) with 5 (35.71%) patients had no pain (VAS=0) in group 1 whereas in the group 2 mean score for leg pain was 0.81 (SD=± 0.89) with no pain in 15 (41.66%) patients (VAS=0). The difference in VAS scores for pain at one week follow up between group 1 and 2 was not significant (p-value=0.682).

3. However, at two weeks follow up the mean leg pain was 0.43 (SD=0.51) in the group 1st while in the group 2nd mean leg pain was 0.11 (SD=0.40). At three weeks follow-up only two patients in group 1 had leg pain while in group 2 no patient reported leg pain, on further follow up all patients in both groups reported no leg pain.

4. There was no significant difference between the mean VAS score for leg pain preoperatively, 24 hours post-operatively, at 1st week, 2nd week, 3rd week, 1st month, 2nd month and 3rd month of follow-up in group 1 and group 2.

Back pain:

1. The pre-operative back pain scores were 6.57 in group 1 and 4.52 in group 2. Scores reduced to 2.29 in group 1 and 1.58 in group 2 24hrs post-operatively. There was a statistically significant difference (p-value=0.019 and 0.008) in the VAS scores for back pain pre-operatively and 24 hrs post-operatively: Patients with upper lumbar level disc prolapse had more back pain than patients with lower lumbar disc prolapse pre-operatively and at initial follow-up at 24 hrs post-operatively.

2. On follow at one week the mean back pain was 0.86 (SD=± 0.95) in group 1 while in the group 2 mean VAS score was 0.44 (SD=± 0.90). The difference in VAS scores for back pain at one week follow up between group 1 and group 2 was NOT significant (p-value=0.554). However, at two weeks follow up the mean back pain was 0.36 (SD=0.63) in the group 1 while in the group 2 mean was 0.17 (SD=0.74). On further follow up patient's in group 1 had some residual mild back pain while patients in group 2 had no back pain except for one patient who had complication of wound infection.

3. There was no significant difference between the mean VAS score for back pain at 1st week, 2nd week, 3rd week, 1st month, 2nd month and 3rd month at follow-up in group 1 and group 2 though there is significant difference pre-operatively and initially postoperatively at 24 hours. Back pain was more significant in upper lumbar disc prolapses. (Table 8).

According to Adam Pearson and et al., Surgery resulted in greater improvement in both back ache, leg pain than non-operative treatment at each follow-up period; however, leg pain improved significantly more than back pain. The treatment effect of surgery was greater for leg pain than back pain at 3 months but not at 1 or 2 years.

Functional status

The disability due to prolapsed intervertebral disc was assessed using the Modified Oswestry Disability Index (MODI) determined before surgical procedure then at 1st month, 2nd month and then at 3rd month after the surgical procedure. The mean ODHI in pre-operative period was 52.29 for group 1 and 50.28 for group 2. At 1st month follow up ODHI in group 1 reduced to 10.43,6.71 at 2nd month and then to 5.43 at 3rd month whereas in group 2 it was 20.94 at 1st month, 12.11 at 2nd month and 6.28 at 3rd month. Thus, initially patients with upper lumbar level disc prolapse showed less disability and better function after discectomy as compared to patients with lower lumbar level disc prolapse but at the end of three months there was no significant difference in the functional outcome in both the groups of patients [16].

Similar to our results J. Lurie and et al. found that the relative advantage for surgery was greater for patients with herniation at higher lumbar levels, with non- operative treatment being less effective in these patients compared with those with herniation at L4-L5 and L5-S1 levels. In contrast, Sanderson et al. reported that patients with L1-L2 or L2-L3 disc prolapse had significantly worse surgical outcome than the ones with L3-L4, L4-L5 and L5-S1 discectomies. The extent of improvement in radicular pain, back pain and function and economic status were found to be 58%, 53% and 33%, respectively in his study. On the other hand, the surgical outcome in the L3-L4 group was favorable and similar to L4-L5 and L5-S1 group. The reason for this might be that most of their patients with L1-L2 and L2-L3 disc herniation in his study had previous lumbar disc surgeries [17].

However, the improvement in the functional disability of the patients was similar in both the groups of patients in long term follow-up in our study. This again ascertains the fact that level of herniation has no significant bearing on the functional outcome of the patient in long term if proper decompression has been achieved by discectomy. Mechanical factors do not influence the functional outcome in patients with prolapsed disc after discectomy.

Conclusion

We also found that upper lumbar disc herniation was more common in elderly adults above 50 years of age while lower lumbar disc prolapse was more common in middle age group 40 to 50 years.

- The relative advantage for surgery was greater for patients with herniation at higher lumbar levels initially compared with those with herniation at L4-L5 and L5-S1 levels.
- Mechanical factors like level of disc herniation do not influence the functional outcome in patients with prolapsed lumbar disc after discectomy.
On comparing the results after discectomy of prolapsed intervertebral disc at different levels in the lumbar spine we found no significant difference in the end result and functional outcome of the patients.

References