Minimally Invasive Multi-Level Posterior Lumbar Interbody Fusion Using a Percutaneously Inserted Spinal Fixation System: Technical Tips, Surgical Outcomes

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Objective: There are technical limitations of multi-level posterior pedicle screw fixation performed by the percutaneous technique. The purpose of this study was to describe the surgical technique and outcome of minimally invasive multi-level posterior lumbar interbody fusion (PLIF) and to determine its efficacy.

Methods: Forty-two patients who underwent mini-open PLIF using the percutaneous screw fixation system were studied. The mean age of the patients was 59.1 (range, 23 to 78 years). Two levels were involved in 32 cases and three levels in 10 cases. The clinical outcome was assessed using the visual analog scale (VAS) and Low Back Outcome Score (LBOS). Achievement of radiological fusion, intra-operative blood loss, the midline surgical scar and procedure related complications were also analyzed.

Results: The mean follow-up period was 25.3 months. The mean LBOS prior to surgery was 34.5, which was improved to 49.1 at the final follow up. The mean pain score (VAS) prior to surgery was 7.5 and it was decreased to 2.9 at the last follow up. The mean estimated blood loss was 238 mL (140-350) for the two level procedures and 387 mL (278-458) for three levels. The midline surgical scar was 6.27 cm for two levels and 8.25 cm for three level procedures. Complications included two cases of asymptomatic medial penetration of the pedicle border. However, there were no signs of neurological deterioration or fusion failure.

Conclusion: Multi-level, minimally invasive PLIF can be performed effectively using the percutaneous transpedicular screw fixation system. It can be an alternative to the traditional open procedures.

Key Words: Posterior lumbar interbody fusion · Percutaneous · Minimally invasive surgery.

INTRODUCTION

Posterior lumbar interbody fusion (PLIF) is a widely performed surgical procedure for the management of pain and spinal instability resulting from various degenerative lumbar spine diseases where the anterior column support is inadequate.12,13 Posterior instrumentation is frequently used to augment interbody fusion, and pedicle screws with rods are commonly employed for this purpose. However, traditional open PLIF for instrument implantation requires a large midline incision and extensive dissection of the paraspinal muscles that can increase the risk for complications both in the short and long term.4,14,15 In contrast, minimally invasive lumbar fusion using percutaneous screw fixation significantly diminishes the risk for complications such as iatrogenic soft-tissue damage and postoperative back pain.14 However, the use of minimally invasive mini-open PLIF using the percutaneous screw fixation system for multi-level degenerative spine disease has not been previously described. The difficulty with rod insertion has been a major technical limitation for the use of percutaneous screw fixation involving multiple levels. The current study describes the surgical technique and feasibility of using the mini-open PLIF procedure for multi-level using percutaneous screw fixation with a vertical axis and detachable screw extender (Fig. 1).

MATERIALS AND METHODS

Patient population
During a period ranging from January 2006 to March 2008,
42 consecutive patients who underwent mini-open PLIF were included in this study. The inclusion criteria were: 1) no previous lumbar surgery, 2) instability accompanying spinal stenosis or spondylolisthesis at more than two levels, 3) followed up for more than 12 months, and 4) no severe osteoporosis (the lowest T-score on DEXA bone mineral density ≥3.0). The study group included 14 men and 28 women. The mean follow-up period was 25.3 months (12 months to 43 months) and the mean age was 59.1 (range, 23 to 78 years). The clinical outcome was assessed using the visual analog scale (VAS) and Low Back Outcome Score (LBOS) (Table 1). In addition, the presence of radiological fusion, intra-operative blood loss, the postoperative midline surgical scar and procedure related complications were evaluated.

**Surgical technique**

All patients underwent surgery in the prone position. Following a midline skin incision, an expandable tubular retractor of appropriate length (X-tube, Medtronic Sofamor Danek, Memphis, TN, USA) was inserted at each level. This was followed by central decompression. On the contralateral side, a foraminal decompression was carefully performed, minimizing the muscle retraction. Not only central decompression but also foraminal decompression could be sufficiently performed through the X-tube, minimizing the damage to facet joint. Next, the percutaneous pedicle screws with a vertical axis and detachable extender (Apollon system®, Solco medical, Korea) were inserted without additional skin incision. After dissection of the subdermal plane, the percutaneous screws were inserted following retraction of the skin (Fig. 2). At this time, in cases where the alignment was not well fitted, the rod insertion may be obstructed. In these cases, using the detachable screw extender of the Apollon system, the screw extender holder bar should be removed (Fig. 3). Then, by adjusting the trajectory of the screw head, the rod is manually controlled for insertion. Thus, the rod could be inserted with ease. After distraction of the instrumented vertebrae, the interbody peek cages filled with local bone chips harvested from the lamina and facet joints were inserted at each level. Finally, the

| Parameter               | Finding                               | Points |
|-------------------------|---------------------------------------|--------|
| Current pain            | 7 to 10 cm VAS                        | 0      |
|                         | 5 to 6 cm VAS                         | 3      |
|                         | 3 to 4 cm VAS                         | 6      |
|                         | 0 to 2 cm VAS                         | 9      |
| Employment              | Unemployed because of back pain       | 0      |
|                         | Part time                             | 3      |
|                         | Full time lighter                     | 6      |
|                         | Full time original                    | 9      |
| Domestic chores         | None                                  | 0      |
| odd jobs                | A few but not many                    | 3      |
|                         | Most or all but more slowly normally  | 6      |
| Sport or active         | None                                  | 0      |
| social activities       | Some but much less than before        | 3      |
|                         | Back to previous level                | 9      |
| Resting                 | Resting more than half the day        | 0      |
|                         | Little rest needed occasional         | 4      |
|                         | No need rest                          | 6      |
| Treatment or consultation| More than once per month              | 0      |
|                         | About once per month                  | 2      |
|                         | Rarely                                | 4      |
|                         | Never                                 | 6      |
| Analgesia               | Several times each day                | 0      |
|                         | Almost every day                      | 2      |
|                         | Occasionally                          | 4      |
|                         | Never                                 | 6      |
| Sex life                | Severely affected impossible          | 0      |
|                         | Moderately affected difficult         | 2      |
|                         | Mildly affected                       | 4      |
|                         | Unaffected                            | 6      |
| Sleeping                | Severely affected impossible          | 0      |
|                         | Moderately affected difficult         | 1      |
|                         | Mildly affected                       | 2      |
|                         | Unaffected                            | 3      |
| Walking                 | Severely affected impossible          | 0      |
|                         | Moderately affected difficult         | 1      |
|                         | Mildly affected                       | 2      |
|                         | Unaffected                            | 3      |
| Sitting                 | Severely affected impossible          | 0      |
|                         | Moderately affected difficult         | 1      |
|                         | Mildly affected                       | 2      |
|                         | Unaffected                            | 3      |
| Travelling              | Severely affected impossible          | 0      |
|                         | Moderately affected difficult         | 1      |
|                         | Mildly affected                       | 2      |
|                         | Unaffected                            | 3      |
| Dressing                | Severely affected impossible          | 0      |
|                         | Moderately affected difficult         | 1      |
|                         | Mildly affected                       | 2      |
|                         | Unaffected                            | 3      |

**Table 1. The low back outcome scale of Greenough and Fraser**

VAS: visual analog scale
screws were compressed into each other to create a lumbar lordosis. In all patients, sitting and walking were allowed two days after surgery. The patients were instructed to wear a thoraco-lumbo-sacral orthosis when out of bed until three months after surgery.

**Statistical analysis**

Statistical analysis, including mean values and standard deviations, was performed using SAS 6.12 (SAS institute, Inc., Cary, NC, USA). Comparisons between different points in time were performed using the paired Student’s t-test. Differences were considered statistically significant at $p<0.05$.

**RESULTS**

Forty-two patients were included in this study. There were no cases to be converted into open surgery due to technical difficulty. The number of levels operated on was two levels in 32 patients and three levels in 10 patients. The mean follow-up period was 25.3 months and all patients were followed for a minimum of 12 months. The mean estimated blood loss was 238 mL (140-350) for two levels and 387 mL (278-458) for three levels. There were no cases required a blood transfusion. The mean LBOS prior to surgery was 32.5, which improved to 53.1 at the last follow-up ($p=0.037$). From a preoperative average of 7.5, the VAS score was decreased significantly to 3.0 at 7 days postoperatively and 2.9 at the final follow-up ($p=0.015$). At a minimum of 12 months follow-up, all cases appeared to have solid fusions as judged by the presence of a trabecular bony bridge and less than 5° on dynamic flexion-extension views. The postoperative midline surgical scars were a mean of 6.27 cm for two levels and 8.25 cm for three levels. Complications included two cases of medial penetration of the pedicle border without neurological deficits and one case with a deep wound infection. However, there were no signs of neurological aggravation or fusion failure at the final follow up. Moreover, there were no neurological sequelae as a result of misplaced pedicle screws (Fig. 4).

**DISCUSSION**

The PLIF has been associated with the improvement of the fusion rate while restoring disc height and maintaining vertebral alignment\(^3\). Since the anterior and middle spinal columns
support 80% of the spinal load, placing the bone graft in this load-bearing position makes it subject to compressive forces that enhance bone fusion as predicted by Wolff’s law. In addition, the vertebral body represents 90% of the osseous surface area and receives a more generous vascular supply than the posterolateral elements, which further improve the fusion potential. However, despite the advantages of the satisfactory fusion rate, PLIF procedure itself has some drawbacks. It is known that postoperative back pain is caused by muscle injury during surgery and is directly related to the operation time and external compression force from the retractor. Due to the massive skin incision, the risk for intraoperative blood loss and postoperative back pain is increased. In addition, for the conventional midline approach for screw fixation for multiple levels, forceful retraction of the paraspinal muscles is required to achieve the proper lateral-to-medial screw trajectory because of coronal plane angle of pedicle. Subsequent prolonged wide retraction may result in denervation of the paraspinal musculature. Moreover, the procedure can potentially injure the medial branches of the dorsal rami at adjacent levels and at the level of fusion; this is because these branches are relatively fixed as they run beneath the fibro-osseous mammilloaccessory ligament. On the contrary, percutaneous pedicle screw fixation permits a safe application while preserving soft tissues without relevant blood loss and persisting sequelae, such as muscular denervation, atrophy, and pain. Percutaneous screws can be inserted in the appropriate direction without forceful retraction of the muscles. Due to the technical problems of rod insertion however, the percutaneous transpedicular screw procedure is usually performed at a single level. With multiple levels, percutaneous screw fixation is difficult to perform. The goal of minimally invasive spinal surgery is to achieve the satisfactory results as comparable open procedure with reduced iatrogenic injury. In the current study, a minimally invasive mini-open PLIF was performed at multiple levels in 42 patients using a vertical axis and detachable screw system; the results were satisfactory. Surgical devices used in percutaneous pedicle screw fixation are classified into two major types depending on the rod insertion method. The first type is the geometrical coordinating device called the ‘Sextant’ (Medtronic) and ‘Apollon’ (Scolot) that facilitates pedicle screw fixation by following the circular margin of the arc during rod insertion. The other device is the open screw head type used for direct insertion of the rod (J&J’s ‘Viper system’ and ‘Path-Finder system’). For the multi-level percutaneous pedicle screwing, the initial alignment of the vertebroplasty needle is essential. Despite the use of a ruler guide, the direction of a needle may be altered during passage through the pedicle. The rod cannot be easily inserted, in case of initial misalignment of screws. However, the vertical axis and detachable screw extender system has made this more feasible. If surgeons become accustomed to the percutaneous screw fixation, the alignment with ruler guide would not be needed. Under surgical observation, the alignment of the vertebroplasty needle should be well adjusted. Following this, with the alignment of the needle consistently maintained a needle at all segments was inserted to the desired depth. However, if the rod insertion is obstructed, the screw extender holder bar should be removed. Misalignment of the screw head leads to the difficulty of rod insertion and the primary cause of this may be the difference in each of its depth and failing to keep them in perfect alignment. Removing the extensor bar offers possibility of controlling the angle of the screw head due to its polychy axial characteristics. It enables controlling the axis of rod insertion which facilitates the process of it additionally, and this is what we supplemented to this content. Then, the screw extender could be detached. Following a re-adjustment of the screw head alignment, the rod could be manually inserted with ease. The technical limitations of multi-level posterior pedicle screw fixation by percutaneous techniques that were present with previous sextant systems have been overcome with this method. Although there are potential benefits with the minimally invasive mini-open procedure, the technique also has some of its own drawbacks and limitations. There is a learning curve that must be achieved before the technical skills are acceptable. However, the minimally invasive mini-open PLIF used for multiple levels can provide many potential advantages compared to the conventional PLIF, although technically demanding.

CONCLUSION

The vertical axis and detachable screw extender system for multi-level interbody fusion makes rod manipulation easier to perform. This technique can reduce the size of the midline skin incision as well as the risks for iatrogenic muscle injury. Accordingly, a prompt recovery and a good clinical outcome can be expected even in cases with multiple levels.

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