Long-Term Clinical and Radiological Outcomes of Minimally Invasive Transforaminal Lumbar Interbody Fusion: 10-Year Follow-up Results

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ABSTRACT

Background: Many studies have reported that minimally invasive transforaminal lumbar interbody fusion (MI-TLIF) provides satisfactory treatment comparable to other fusion methods. However, in the case of MI-TLIF, there are concerns about the long-term outcome compared to conventional bilateral PLIF due to the small amount of disc removal and the lack of autogenous bone graft. Long-term follow-up studies are still lacking as most of the previous reports have follow-up periods of up to 5 years.

Methods: Thirty patients who underwent MI-TLIF were followed up for > 10 years (mean, 11.1 years). Interbody fusion rates were determined using a modified Bridwell grading system. Adjacent segment disease (ASD) was defined as radiological adjacent segment degeneration (R-ASDeg) as seen on plain X-rays; reoperated adjacent segment disease referred to the subsequent need for revision surgery. Clinical outcomes after surgery were assessed based on back and leg pain as well as the Oswestry disability index (ODI).

Results: The overall radiological fusion rate, at the 1-, 5-, and 10-year follow-up was 77.1%, 91.4%, and 94.3%, respectively. The incidence of R-ASDeg 1, 5, and 10 years after surgery was 6.7%, 16.7%, and 43.3% at the proximal adjacent segment and 4.8%, 14.3%, and 28.6% at the distal adjacent segment, respectively. R-ASDeg at either the proximal or distal segment was determined in 50.0% of the patients 10 years postoperatively. All clinical parameters improved significantly during follow-up, although the ODI and the visual analog scale (VAS) for leg pain at the 10-year follow-up were significantly worse in the R-ASDeg group than in the other patients (P = 0.009, P = 0.040).

Conclusion: MI-TLIF improved both clinical and radiological outcomes, and the improvements were maintained for up to 10 years after surgery. However, R-ASDeg developed in up to 50% of the patients within 10 years, and both leg pain on the VAS and the ODI were worse in patients with R-ASDeg.

Keywords: Degenerative Lumbar Disease; Long-term Follow Up; Minimally Invasive Transforaminal Interbody Fusion; Ten Years
INTRODUCTION

Lumbar interbody fusion is used to treat many diseases of the lumbar spine, such as spinal stenosis, spondylolisthesis, and spinal deformation. Among the various fusion methods, posterior approaches are most commonly used, including conventional posterior lumbar interbody fusion (PLIF) and transfemoral lumbar interbody fusion (TLIF), first introduced by Harms et al.

Unlike conventional PLIF, TLIF can achieve decompression and interbody arthrodesis by preserving the opposite facet and lamina in a unilateral approach, while reducing the amount of soft-tissue injury as well as the extent of thecal-sac retraction. Nevertheless, both muscle atrophy due to the extensive skin incisions and perivertebral muscle detachment are common postoperative problems. Recently, minimally invasive techniques using a tubular retractor have been shown to preserve much of the lumbar musculature while splitting the muscle to greatly reduce intraoperative bleeding. Among the advantages are a reduction in pain immediately after surgery, a faster recovery, and a shorter hospital stay and thus a faster return to daily life.

Many studies have reported that minimally invasive TLIF (MI-TLIF) provides satisfactory treatment comparable to other fusion methods, with outcomes similar to those obtained with the conventional fusion method in terms of interbody fusion, complications, and clinical results. However, in MI-TLIF annulectomy and disc resection are performed on only 1 side and only a single cage is inserted, such that the area of disc space preparation may be smaller than in a conventional bilateral PLIF. Furthermore, since the laminar resection is unilateral, much less autogenous local bone can be harvested and an allograft is thus additionally required. These issues have raised concern as to whether, over the long term, the fusion rate is lower than that achieved with conventional PLIF. Nonetheless, an advantage of MI-TLIF is that it minimizes the impact on the connection between the index level and the adjacent segment by preserving the surrounding structures and the interconnecting soft tissues, especially on the contralateral side. Accordingly, compared to conventional open interbody fusion, MI-TLIF should be less likely to cause adjacent segment degeneration (ASD). However, long-term follow-up studies are lacking as most reports have a maximum follow-up of 5 years.

Here we report the clinical and radiological results obtained in a 10-year follow-up of patients with degenerative lumbar disease treated by MI-TLIF.

METHODS

Study design and patient selection
A retrospective review was conducted of patients who, between September 2005 and August 2010, underwent MI-TLIF followed by percutaneous pedicle screw fixation (PPSF) at a single institution. Adult (≥18 years of age) patients with a radiological and clinical follow-up of at least 10 years after MI-TLIF for spinal stenosis, spondylolisthesis, degenerative disc disease, or other degenerative lumbar diseases involving L3 to S1 were included in the study. Patients with a previous history of infection, trauma, tumor, or previous surgery on the operated or adjacent segment, diseases and drugs that affect bone metabolism, or incomplete follow-up data were excluded. Among the 107 consecutive patients who underwent MI-TLIF, 86 patients...
were followed for 1 year or more, 65 patients for 3 years or more, 42 patients for 5 years or more, and 30 patients for 10 years or more. The follow-up rates are 80.4%, 60.7%, 39.3%, and 28.0%, respectively, showing a gradually decreasing pattern as time goes by. In this study, 30 patients (28.0%) who had been followed-up for at least 10 years were included.

**Surgical procedures**

The patients were administered general anesthesia and placed in a prone position. A 22-mm longitudinal incision was then made with a disc space trajectory on the symptomatic side and a dilater was inserted continuously under fluoroscopic guidance, followed by the insertion of a tubular retractor system (METRx; Medtronic Sofamor Danek, Memphis, TN, USA). Only the facet joint on the symptomatic side was removed, using a high-speed burr with an L-shaped osteotome. After removal of the thickened yellow ligament to decompress the dural sac and nerve roots, the end plate was prepared and the patient’s local bone, collected during surgery, was implanted into the disc space. Additional allogeneic bone was used as needed. The cage (either Capstone [Medtronic Sofamor Danek, Minneapolis, MN, USA], or OIC [Stryker Spine, Allendale, NJ, USA]) was filled solely with autogenous bone chips and then sealed with demineralized bone matrix (DBM, Orthoblast II; Isotis Orthobiologics, Irvine, CA, USA). The prepared cage was then placed on the more symptomatic side. Contralateral nerve decompression, when required, was performed by changing the angle of the tubular retractor and using unilateral laminotomy and bilateral decompression methods. After decompression and fusion were completed, the tubular retractor was removed and PPSF was performed. A detailed description of the procedure was provided in a previous paper.21

**Measurement of radiological outcomes**

Radiological parameters were evaluated based on the preoperative and 1-, 5-, and 10-year postoperative follow-up examinations. All parameters were measured using picture archiving and communication system (PACS) software and PACS workstations. Disc height and foraminal height were measured on a plain lateral radiograph together with segmental lordosis of the operated segment and lumbar lordosis.

Interbody fusion rates were assessed with the modified Bridwell interbody fusion grading system using both plain radiograph and computed tomography imaging (grade 1: fused with bony bridging, presence of trabecular remodeling; grade 2: not fully bony bridged, remodeling but without radiolucency above or below the cage; grade 3: radiolucency present at the top or bottom of the cage and screw; grade 4: fusion absent, with false motion). In this study, Bridwell grades 1 and 2 were defined as fused and Bridwell grades 3 and 4 as non-fused. Interobserver agreement of the modified Bridwell grade was assessed by 2 independent spinal surgeons. The kappa value was 0.78, indicating substantial agreement according to the Landis and Koch criteria.22

Disc degeneration of the adjacent segments was evaluated preoperatively using magnetic resonance (MR) imaging according to Pfirrmann’s classification.23 Preoperative stenosis of the spinal canal was evaluated using T2-weighted sagittal MR images according to Imagama’s classification.24

Radiological adjacent segment degeneration (R-ASDeg) was evaluated on plain X-rays, with separate assessments of the proximal and distal segments.25 R-ASDeg positivity was diagnosed based on comparisons with the preoperative radiographs and defined as a reduction in disc height by ≥ 3 mm on a neutral lateral radiograph, an increase in vertebral

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slip of ≥ 3 mm as seen on a neutral lateral radiograph, and an increase in the posterior opening angle of ≥ 5° as seen on a flexion lateral radiograph.

**Measurement of clinical outcomes**
The demographic information collected in this study included age, sex, body mass index, smoking status, osteoporosis, and disease history. Data on the patients' surgical parameters, including the preoperative diagnosis, operation time, intraoperative blood loss, number of operated levels, operated level, and hospitalization period, were obtained from the medical records.

Clinical outcomes were assessed using the visual analog scale (VAS) for back pain and leg pain and the Oswestry disability index (ODI). All clinical variables were measured preoperatively and at 1, 5, and 10 years postoperatively. Perioperative complications, such as wound infection, postoperative neuralgia, and dural tear, were recorded, as well as the occurrence of pseudarthrosis, implant breakage, screw loosening, and cage subsidence as late complications. Reoperated adjacent segment disease (O-ASD) described those cases in which additional surgery was required to treat pain or neurological symptoms resulting from ASD.

Clinical results, including the VAS score for back pain and leg pain and the ODI, were compared between patients who underwent 1-level and 2-level surgery. The differences in clinical outcomes of young (< 60 years) vs. older (≥ 60 years) MI-TLIF-treated patients was also determined.

**Statistical analysis**
Clinical and radiological outcomes were analyzed using Student's t-test and the Mann-Whitney test for continuous variables, and a χ² test or Fisher's exact test for categorical variables. Differences in the clinical and radiological outcomes over time as described by the continuous variables were analyzed using a repeat measures analysis of variance. A generalized estimation equation was used to compare the radiological and clinical outcomes between groups based on the presence or absence of R-ASDeg. The survival rates of R-ASDeg and O-ASD were analyzed according to follow-up period and fusion length using the Kaplan-Meier method. The survival rates of R-ASDeg and O-ASD were analyzed according to follow-up period and fusion length using the Kaplan-Meier method. The statistical analyses were conducted using SPSS software (IBM SPSS statistics for Windows, Version 25.0; IBM, Armonk, NY, USA). A P value < 0.050 was considered to indicate statistical significance.

**Ethics statement**
The present study protocol was reviewed and approved by the Institutional Review Board of Soonchunhyang University Seoul Hospital (approval No. SCHUH 2021-01-008). Informed consent was waived because of the retrospective nature of the study.

**RESULTS**
Thirty patients underwent MI-TLIF and were followed-up for at least 10 years. The mean duration of follow-up was 11.1 years (range, 10.0–12.7 years). The mean age at surgery was 60.1 ± 11.3 years (range, 40–80 years). The number of fused levels was 1 segment in 25 patients (83.3%), and 2 segments in 5 patients (16.7%). L4–5 was the most frequently operated level, followed by L5–S1 and L3–4. The main pathological indications for surgery were spinal stenosis (n = 14, 46.7%), spondylolisthesis (n = 13, 43.3%), recurrent
disc herniation (n = 2, 6.7%), and internal disc disruption (n = 1, 3.3%). The baseline characteristics of the patients are listed in Table 1.

**Radiological outcomes**

A significant increase in foraminal height compared to the preoperative value was determined 1 year postoperatively and was maintained at the 10-year follow-up (P < 0.001). Slight increases in mean disc height, segmental lordosis, and lumbar lordosis were measured 1 year, 5 years, and 10 years postoperatively, but the difference compared to the initial value was not significant (Table 2).

Interbody fusion was evaluated using a modified Bridwell grade and differed significantly between the 1-year and the 5-year follow-up (P < 0.001) and between the 1-year and 10-year follow-up (P < 0.001), but not between the 5-year and 10-year follow-up. The ratio of Bridwell grades 1 and 2 (fused) increased gradually, from 27 segments (77.1%) at 1 year to 32 segments (91.4%) at 5 years, and 33 segments (94.3%) at 10 years postoperatively. The difference in the fusion rate 1 and 10 years postoperatively was significant (P = 0.041) (Fig. 1). Among our patients, 2 had pseudarthrosis, which persisted until the last follow-up but remained asymptomatic and was managed conservatively.

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**Table 1. Patient demographic and operative data**

| Variables                      | Values          |
|--------------------------------|-----------------|
| Age at surgery, yr            | 60.1 ± 11.3     |
| Sex, M:F                       | 9:21            |
| BMI, kg/m²                     | 26.3 ± 3.8      |
| Smoker                         | 2 (6.7)         |
| Diabetes                       | 12 (40.0)       |
| Hypertension                   | 14 (46.7)       |
| Osteoporosis                   | 14 (46.7)       |
| Surgery level                  |                 |
| Single level                   | 25 (83.3)       |
| Two levels                     | 5 (16.7)        |
| Mean number of levels fused    | 1.2 ± 0.4       |
| Preoperative diagnosis         |                 |
| Spondylolisthesis              | 13 (43.3)       |
| Spinal stenosis                | 14 (46.7)       |
| Recurred disc herniation       | 2 (6.7)         |
| Internal disc disruption       | 1 (3.3)         |
| Treated segments               |                 |
| L3–4                           | 6 (17.1)        |
| L4–5                           | 20 (57.1)       |
| L5–S1                          | 9 (25.7)        |
| Follow-up duration, yr         | 11.1 ± 0.8      |
| Operative time per segment, min| 201.2 ± 52.3    |
| Average intraoperative blood loss, mL | 213.3 ± 126.4 |
| Hospital stay, day             | 22.6 ± 24.3     |

Data are presented as mean ± standard deviation or number (%). BMI = body mass index.

**Table 2. Radiologic outcomes**

| Variables                      | Initial | 1 yr | 5 yr | 10 yr | P value |
|--------------------------------|---------|------|------|-------|---------|
| Disc height, mm                | 8.5 ± 1.4| 9.0 ± 1.2 | 9.2 ± 1.7 | 9.1 ± 1.7 | 0.172 |
| Foraminal height, mm           | 16.2 ± 1.8| 17.8 ± 2.2 | 17.6 ± 2.1 | 17.5 ± 2.1 | < 0.001*** |
| Segmental lordosis, °          | 14.0 ± 7.7| 14.6 ± 5.5 | 14.0 ± 5.1 | 14.2 ± 5.6 | 0.658 |
| Lumbar lordosis, °             | 40.4 ± 12.2| 42.7 ± 9.4 | 41.3 ± 9.5 | 41.7 ± 9.7 | 0.178 |

Data are presented as mean ± standard deviation. ***P < 0.001.
R-ASDeg occurred more frequently in the proximal than in the distal adjacent segment, but the difference was not significant. A reduction in disc height of > 3 mm, as determined at the 10-year follow-up, more often involved the proximal (6/30, 20.0%) than the distal (3/21, 14.3%) adjacent segment. The incidence of an increase of the posterior opening angle at the adjacent segment of > 5° did not significantly differ between the proximal (2/30, 6.7%) and distal (1/21, 4.8%) adjacent segments at 10 years, nor did the incidence of an increase in vertebral slip of > 3 mm (proximal adjacent segment: 3/30, 10.0%; distal adjacent segment: 2/21, 9.5%). The incidence of any of these R-ASDeg signs (disc height reduction, increased posterior opening angle, and increased vertebral slip) at the 1-year, 5-year, and 10-year follow-up was 6.7%, 16.7%, and 43.3% at the proximal adjacent segment, and 4.8%, 14.3%, and 28.6% at the distal adjacent segment, respectively. Despite a tendency of a greater frequency in the proximal than in the distal segment, the difference was not significant. A gradual increase in R-ASDeg involving either the proximal or the distal adjacent segment was determined, with rates of 10.0%, 26.7%, and 50.0% 1, 5, and 10 years postoperatively, but the differences between the corresponding values of the proximal and distal adjacent segments were not significant (Table 3).

Table 3. The R-ASDeg incidence at the proximal and distal adjacent levels

| Variables                              | 1 yr | 5 yr | 10 yr | P value |
|----------------------------------------|------|------|-------|---------|
| The incidence of disc height reduction (≥ 3 mm) |      |      |       | 0.587   |
| Proximal (n = 30)                      | 1 (3.3) | 2 (6.7) | 6 (20.0) |         |
| Distal (n = 21)                        | 1 (4.8) | 2 (9.5) | 3 (14.3) |         |
| P value                                | 1.000 | 1.000 | 0.720 |         |
| The incidence of an increase of posterior opening angle (≥ 5°) |      |      |       | 0.320   |
| Proximal (n = 30)                      | 1 (3.3) | 2 (6.7) | 2 (6.7)   |         |
| Distal (n = 21)                        | 0 (0.0) | 0 (0.0) | 1 (4.8) |         |
| P value                                | 1.000 | 0.506 | 1.000 |         |
| The incidence of vertebral slip (an increase of ≥ 3 mm) |      |      |       | 0.962   |
| Proximal (n = 30)                      | 0 (0.0) | 1 (3.3) | 3 (10.0) |         |
| Distal (n = 21)                        | 0 (0.0) | 1 (4.8) | 2 (9.5) |         |
| P value                                | -     | 1.000 | 1.000 |         |
| The total incidence of R-ASDeg          |      |      |       | 0.706   |
| Proximal (n = 30)                      | 2 (6.7) | 5 (16.7) | 10 (41.3) |         |
| Distal (n = 21)                        | 1 (4.8) | 3 (14.3) | 6 (28.6) |         |
| P value                                | 1.000 | 1.000 | 0.718 |         |

Data are presented as number (%).
R-ASDeg = radiological adjacent segment degeneration.
In the analysis of patients classified according to the presence or absence of R-ASDeg, in the former the VAS score for leg pain was significantly higher at the 10-year follow-up \( (P = 0.040) \), while the ODI was significantly higher both at 5 years \( (P = 0.002) \) and 10 years \( (P = 0.009) \) (Fig. 2). The 2 groups did not significantly differ with respect to back pain and other variables (Table 4).

**Clinical outcomes**

All clinical parameters improved significantly after surgery and the improvements were maintained over the 10-year follow-up period. Both the VAS score for back and leg pain and the ODI improved significantly at the 1-year, 5-year, and 10-year follow-up examinations compared to their preoperative values, without a significant difference over time (Fig. 3). The clinical scores were further analyzed by dividing patients into those with 1-level vs. 2-level surgery. The VAS score and ODI improved in each group but the changes over time were similar in the 2 groups (Table 5). MI-TLIF was as effective in patients with a 2-level pathology as in those with a 1-level pathology. There was also no difference in the clinical scores between older and young patients based on a cut-off of 60 years of age (Supplementary Table 1).

No deaths or serious complications occurred. One patient developed a surgical site infection soon after surgery but it was successfully treated by superficial incision and drainage and resolved by 2 weeks after MI-TLIF. Two patients complained of postoperative neuralgia at the first outpatient visit, but both responded well to pregabaline. Late complications consisted of mild cage subsidence of < 25% of the disc height in 9 patients (30.0%); however, the subsidence did not cause neurological symptoms and secondary surgery was not required. As interbody fusion progressed, the subsidence ceased in all patients. A screw halo around the screw, visible on the plain anteroposterior radiograph and computed tomography images, occurred in 2 patients (6.7%), but as the union progressed the halo filled with bone and, at the last follow-up, solid interbody union was observed.

**Incidence of secondary surgery**

Seven of the 30 patients required reoperation, as determined at the last follow-up. The mean time from the first operation to reoperation was 7.7 years (range, 3.5–12.2). Among these
patients, O-ASD involved the proximal adjacent segments in 3 patients, the distal adjacent segments in 3 patients, and both the proximal and the distal adjacent segments in 1 patient.

The pathology leading to O-ASD was spinal stenosis in 5 patients, degenerative listhesis in 1 patient, and intervertebral disc herniation in 1 patient. Reoperation consisted of open PLIF, with spinal fusion extended to adjacent segments in the 6 patients with spinal stenosis or degenerative listhesis. Microdiscectomy was performed in the patients with disc herniation. In all 7 O-ASD patients, both the neurological symptoms and pain improved after the second surgery.

Table 4. Comparison of patients with and without R-ASDeg at 10 years after primary surgery

| Variables                        | Without R-ASDeg (n = 15) | With R-ASDeg (n = 15) | P value |
|----------------------------------|--------------------------|-----------------------|---------|
| Age, yr                          | 58.7 ± 9.3               | 61.5 ± 13.2           | 1.000   |
| Sex, M:F                         | 3:12                     | 6:9                   | 0.427   |
| BMI, kg/m²                       | 26.5 ± 4.4               | 26.1 ± 3.2            | 0.799   |
| Smoker                           | 1 (6.7)                  | 1 (6.7)               | 1.000   |
| Diabetes                         | 7 (46.7)                 | 5 (33.3)              | 0.456   |
| Hypertension                     | 5 (33.3)                 | 9 (60.0)              | 0.272   |
| Osteoporosis                     | 7 (46.7)                 | 7 (46.7)              | 1.000   |
| Mean number of levels fused      | 1.3 ± 0.5                | 1.1 ± 0.3             | 0.155   |
| Preoperative diagnosis           |                          |                       | 0.355   |
| Spondylolisthesis                | 6 (40.0)                 | 7 (46.7)              |         |
| Spinal stenosis                  | 8 (53.3)                 | 6 (40.0)              |         |
| Recurrent disc herniation        | 0 (0.0)                  | 2 (13.3)              |         |
| Internal disc rupture            | 1 (6.7)                  | 0 (0.0)               |         |
| Back pain                        |                          |                       |         |
| 1 yr after surgery               | 1.8 ± 1.4                | 2.7 ± 2.5             | 0.259   |
| 5 yr after surgery               | 1.5 ± 1.5                | 2.1 ± 2.0             | 0.353   |
| 10 yr after surgery              | 2.2 ± 2.1                | 2.8 ± 2.0             | 0.439   |
| Leg pain                         |                          |                       |         |
| 1 yr after surgery               | 1.3 ± 2.2                | 1.8 ± 3.1             | 0.585   |
| 5 yr after surgery               | 0.7 ± 1.4                | 2.0 ± 2.1             | 0.663   |
| 10 yr after surgery              | 1.0 ± 1.6                | 2.3 ± 1.7             | 0.040*  |
| ODI score                        |                          |                       |         |
| 1 yr after surgery               | 9.9 ± 8.4                | 12.5 ± 9.7            | 0.452   |
| 5 yr after surgery               | 6.7 ± 4.6                | 14.5 ± 7.2            | 0.002** |
| 10 yr after surgery              | 6.5 ± 13.3               | 13.3 ± 7.8            | 0.009** |
| Adjacent segment disc height     |                          |                       |         |
| Proximal (n = 30)                | 10.2 ± 1.9               | 11.0 ± 3.3            | 0.419   |
| Distal (n = 21)                  | 12.0 ± 1.2               | 13.7 ± 3.2            | 0.167   |
| Adjacent segment body listhesis, mm |                      |                       |         |
| Proximal (n = 30)                | 2.1 ± 1.2                | 2.0 ± 1.6             | 0.895   |
| Distal (n = 21)                  | 2.3 ± 1.0                | 2.5 ± 1.6             | 0.650   |
| Adjacent segment intervertebral angle in flexion, ° |          |                       |         |
| Proximal (n = 30)                | 3.1 ± 3.2                | 4.6 ± 3.5             | 0.250   |
| Distal (n = 21)                  | 5.5 ± 3.1                | 6.2 ± 3.9             | 0.693   |
| Adjacent segment preoperative disc degeneration grade |       |                       |         |
| Proximal (n = 30)                | 3.4 ± 0.7                | 3.1 ± 1.1             | 0.338   |
| Distal (n = 21)                  | 3.3 ± 0.9                | 2.7 ± 0.8             | 0.139   |
| Adjacent segment preoperative SCS grade |                      |                       |         |
| Proximal (n = 30)                | 2.3 ± 0.7                | 2.5 ± 0.8             | 0.489   |
| Distal (n = 21)                  | 2.4 ± 0.5                | 2.2 ± 0.6             | 0.580   |

Data are presented as mean ± standard deviation or number (%).

R-ASDeg = radiological adjacent segment degeneration, BMI = body mass index, ODI = Oswestry disability index, SCS = spinal canal stenosis.

*P < 0.05; **P < 0.01.
Survival analysis was performed by dividing the patients into two groups, those with less than 10 years of follow-up and those with more than 10 years of follow-up. No significant difference was observed in the R-ASDeg and O-ASD survival curves between the two groups (R-ASDeg, \( P = 0.889 \); O-ASD, \( P = 0.946 \)). No significant difference was observed in the R-ASDeg and O-ASD survival curves between the two groups even when survival analysis was performed for patients who underwent 1-level fusion surgery and those who underwent 2-level fusion surgery (R-ASDeg, \( P = 0.349 \); O-ASD, \( P = 0.999 \)) (Fig. 4).

DISCUSSION

As life expectancy in many countries continues to increase, due to advances in medical technology, the demand for effective spinal surgery is also likely to increase. Previously, most studies examined perioperative complications, but the focus of recent research has shifted to a consideration of the long-term results, especially long-term clinical outcomes and the inevitable occurrence of ASD following fusion surgery. The goal of minimally invasive techniques is to preserve the paraspinal muscles and soft tissues and thus maintain spinal...
stability between the index and adjacent segments. For this reason, MI-TLIF has been widely adopted and its results are comparable to those achieved with conventional PLIF or open TLIF. However, most of the studied patients have been followed-up for < 5 years. In our long-term study, while the union rate 1 year after MI-TLIF was relatively low (77%), it increased to 91% at 5 years and 94% at 10 years, thus ultimately resulting in a high rate of union (Fig. 5). A limitation of MI-TLIF compared to conventional open PLIF is that the volume of local bone available as a graft is limited, because the procedure includes only a hemilaminectomy, performed through a tubular retractor. Although an allo-chip bone graft and DBM are used to compensate for this deficit, there is concern that it will lead to a low rate of union. Bone morphogenic protein (BMP) was not used in any of our patients. Nonetheless, our study was able to demonstrate the reliability of MI-TLIF as a fusion method as well as successful spinal fusion even without BMP, with results comparable to those reported in previous studies. For example, in a study that included MI-TLIF patients with a minimum 5-year follow-up, the overall fusion rate was 97.7%, with complete fusion and interbody bony

Fig. 4. Kaplan-Meier survival curves. (A) R-ASDeg according to follow-up period. (B) O-ASD according to follow-up period. (C) R-ASDeg according to fusion length. (D) O-ASD according to fusion length. R-ASDeg = radiological adjacent segment degeneration, O-ASD = reoperated adjacent segment disease.
bridging achieved in 84.1%. In another study of MI-TLIF in which patients were followed-up for 5 years, solid fusion was achieved in 72% of the patients by the second postoperative year and in 74% by the fifth postoperative year. Bin Abd Razak et al. performed 1-level MI-TLIF in patients with spondylolisthesis and reported grade 1 fusion 6 months, 2 years, and 5 years postoperatively in 39.3%, 92.9%, and 96.4%, respectively. However, in their study, in 15 patients the bone graft included additional autogenous bone from the posterior iliac crest, and in 41 patients a larger amount of DBM was used than in our study.

Several studies have described ASD after lumbar fusion surgery, but the incidence varied depending on the definition of ASD. The reported incidence of R-ASDeg is 31–82.6% whereas the incidence of O-ASD is much lower, 4.9–24%. The incidence of ASD also varies because it depends on the duration of follow-up. In our study, at the 10-year follow-up exam the incidence of R-ASDeg was 50% and that of O-ASD 23%. When these results are compared with those of previous studies of open PLIF, in which the patients were followed-up for at least 10 years, the incidence of R-ASDeg was lower in our study. Okuda et al. reported that R-ASDeg developed in 75% and O-ASD in 15% of patients with degenerative spondylolisthesis who underwent open PLIF and were followed-up for > 10 years. In a similar study, Nakashima et al. reported R-ASDeg in 68% and O-ASD in 9.9%. The lower incidence of O-ASD than in our study may have been due to differences in the eligibility criteria for reoperation. Further prospective studies are needed to determine whether the 2 surgical methods result in different rates of ASD.

The R-ASDeg observed in our study tended to involve the proximal adjacent segments more often than the distal adjacent segments, as also described by other investigators. In Bydon et al., disc degeneration and reoperation in patients with ASD who underwent L4–5 discectomy were seen more often in the proximal than in the distal adjacent segments. Another study reported that 24 patients who underwent L4–5 fusion surgery required reoperation due to ASD of the proximal adjacent segment, L3–4, compared to 3 patients reoperated for ASD of the distal adjacent segment. Among our patients, L4–5 was the most frequently operated site. The distal adjacent segment, L5–S1, is supported by the tightly connected lumbosacral ligaments, which limits abnormal movement, and by the sacrum and pelvis, which increase biomechanical stability.
However, the implications of R-ASDeg are controversial, with at least 2 studies failing to find an association of R-ASDeg with clinical symptoms. While the incidence of R-ASDeg and O-ASD differ significantly, our patients with R-ASDeg suffered statistically worse leg pain, as evaluated using the VAS and ODI. This result suggests that R-ASDeg is related to the deterioration of clinical symptoms and is a pre-stage to O-ASD progression.

Several studies have reported long-term clinical results after MI-TLIF. In the series of Bin Abd Razak et al., the improvements in the VAS for back and leg pain and in the ODI, 6 months after surgery, were maintained for up to 5 years; similar results were obtained with the Short-Form 36 health survey and Neurogenic Symptoms Score. In the study of Rouben et al., improvements in the ODI and in the VAS of back pain, determined 3 months postoperatively, were maintained at 49 months, regardless of the preoperative diagnosis. In our study, the clinical outcomes after MI-TLIF were maintained for a longer period of time than in previous studies; in our patients the improvements in back pain and leg pain in the first postoperative year were maintained 10 years after surgery. In other MI-TLIF studies the follow-up time was limited to 5 years, whereas our study shows that MI-TLIF continues to provide good clinical results even 10 years after the procedure.

The analyses of our MI-TLIF patients, classified according to the fusion level length, preoperative diagnosis, and age showed that, similar to other studies, these subgroups did not differ in terms of the postoperative improvements in back pain and leg pain on VAS and ODI, either after surgery or at the 10-year follow-up. Although a previous study in which patients were assessed using the VAS and ODI reported greater improvement in those with multi-level than with 1-level fusion, the preoperative VAS and ODI were higher in the 1-level group whereas the postoperative values were similar to those of the latter; thus, the trend over time was in agreement with our findings. In another study, patients treated with 1-level MI-TLIF were classified into isthmic spondylolisthesis and degenerative spondylolisthesis subgroups based on the preoperative diagnosis. Outcomes were evaluated using the VAS and ODI as well as patient satisfaction and return to work rates, but none of the differences between the subgroups were significant. These results together with those of other studies demonstrate the good long-term clinical outcomes that can be achieved with MI-TLIF, regardless of the preoperative diagnosis, patient age, or the need for 2-stage fusion.

Failure to follow-up in long-term follow-up studies can lead to misinterpretation and biased conclusions. The possibility that follow-up rates will not be the same between patients with good and bad clinical outcomes is always a difficult question to answer and a common limitation of human studies. Interestingly, in a recent study comparing the clinical outcomes of patients with and without postoperative follow-up for lumbar degenerative disease, there were no significant differences between groups. In this study, to minimize this limitation, the survival rate was analyzed by dividing the group with less than 10 years of follow-up and the group with more than 10 years of follow-up using the Kaplan-Meier method. When survival analysis was conducted using R-ASDeg and O-ASD as events, there was no significant difference in the survival rate between groups according to follow-up period. Therefore, although the follow-up rate of this study is not high, it is expected to provide meaningful results and will be a valuable reference for future randomized prospective studies.

A strength of our study was that it was performed at a single center and all of the operations were performed by a single senior spine surgeon (Jae Chul Lee). These features increased the reliability of the study, by reducing the number of variables. Another advantage of our study...
was a follow-up of > 10 years after MI-TLIF. However there were also limitations that should be addressed. These included the retrospective design, which may have resulted in observer bias, a lack of standardization of the data, and a lower quality than prospective studies, as well as the relatively small population. This resulted in a lack of statistical power in some of the analyses. Despite these weaknesses, our study is the only one thus far to present 10-year follow-up data for MI-TLIF. Studies of patients followed prospectively for at least 10 years will provide further insights into the clinical advantages of MI-TLIF.

In conclusion, in patients with lumbar spine disease treated by MI-TLIF, the improved clinical and radiological outcomes obtained postoperatively were maintained for up to 10 years and the fusion rates at the end of follow-up were high. However, during the first 10 postoperative years, 50% of patients developed R-ASDeg, which was also characterized by worse leg pain as assessed using the VAS and ODI tools.

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The English in this document has been checked by at least two professional editors, both native speakers of English. For a certificate, please see: http://www.textcheck.com/certificate/FiagTh.

SUPPLEMENTARY MATERIAL

Supplementary Table 1
Young vs. elderly (≥ 60) clinical outcomes

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REFERENCES

1. Lee BH, Moon SH, Suk KS, Kim HS, Yang JH, Lee HM. Lumbar spinal stenosis: pathophysiology and treatment principle: a narrative review. Asian Spine J 2020;14(5):682-93.
2. Park Y, Ha JW. Comparison of one-level posterior lumbar interbody fusion performed with a minimally invasive approach or a traditional open approach. Spine (Phila Pa 1976) 2007;32(5):537-43.
3. Ntoukas V, Müller A. Minimally invasive approach versus traditional open approach for one level posterior lumbar interbody fusion. Minim Invasive Neurosurg 2010;53(1):21-4.
4. Kim YH, Ha KY, Rhyu KW, Park HY, Cho CH, Kim HC, et al. Lumbar interbody fusion: techniques, pearls and pitfalls. Asian Spine J 2020;14(5):730-41.
5. Harms J, Jeszenszky D. The unilateral transformaminal approach for posterior lumbar interbody fusion. Orthop Traumatol 1998;6:88-99.
6. Humphreys SC, Hodges SD, Patwardhan AG, Eck JC, Murphy RB, Covington LA. Comparison of posterior and transformaminal approaches to lumbar interbody fusion. Spine (Phila Pa 1976) 2001;26(5):567-71.
7. Foley KT, Holly LT, Schwender JD. Minimally invasive lumbar fusion. Spine (Phila Pa 1976) 2003;28(15 Suppl):S26-35.
8. Hee HT, Castro FP Jr, Majd ME, Holt RT, Myers L. Anterior/posterior lumbar fusion versus transforaminal lumbar interbody fusion: analysis of complications and predictive factors. *J Spinal Disord* 2001;14(6):533-40.

9. Brodke DS, Dick IC, Kunz DN, McCabe R, Zdeblick TA. Posterior lumbar interbody fusion. A biomechanical comparison, including a new threaded cage. *Spine (Phila Pa 1976)* 1997;22(1):26-31.

10. Miyakoshi N, Abe E, Shimada Y, Okuyama K, Suzuki T, Sato K. Outcome of one-level posterior lumbar interbody fusion for spondylolisthesis and postoperative intervertebral disc degeneration adjacent to the fusion. *Spine (Phila Pa 1976)* 2000;25(14):1837-42.

11. Ringel F, Stoffel M, Stüer C, Meyer B. Minimally invasive transmuscular pedicle screw fixation of the thoracic and lumbar spine. *Neurosurgery* 2006;59(4 Suppl 2):ONS361-6.

12. Schwender JD, Holly LT, Rouben DP, Foley KT. Minimally invasive transforaminal lumbar interbody fusion (TLIF): technical feasibility and initial results. *J Spinal Disord Tech* 2005;18 Suppl:S1-6.

13. Park J, Ham DW, Kwon BT, Park SM, Kim HJ, Yeom JS. Minimally invasive spine surgery: techniques, technologies, and indications. *Asian Spine J* 2020;14(5):694-701.

14. Peng CW, Yue WM, Poh SY, Yeo W, Tan SB. Clinical and radiological outcomes of minimally invasive versus open transforaminal lumbar interbody fusion. *Spine (Phila Pa 1976)* 2009;34(13):1385-9.

15. Scheufler KM, Dohmen H, Vougioukas VI. Percutaneous transforaminal lumbar interbody fusion for the treatment of degenerative lumbar instability. *Neurosurgery* 2007;60(4 Suppl 2):203-12.

16. Schizas C, Tsiridis E, Kosmopoulos V. Minimally invasive versus open transforaminal lumbar interbody fusion: evaluating initial experience. *Int Orthop* 2009;33(6):1683-8.

17. Chang HK, Huang M, Wu JC, Huang WC, Wang MY. Less opioid consumption with enhanced recovery after surgery transforaminal lumbar interbody fusion (TLIF): a comparison to standard minimally-invasive TLIF. *Neurosurg Focus* 2020;17(2):228-36.

18. Park P, Foley KT. Minimally invasive transforaminal lumbar interbody fusion with reduction of spondylolisthesis: technique and outcomes after a minimum of 2 years' follow-up. *Neurosurg Focus* 2008;25(2):E16.

19. Park Y, Ha JW, Lee YT, Sung NY. The effect of a radiographic solid fusion on clinical outcomes after minimally invasive transforaminal lumbar interbody fusion. *Spine J* 2011;11(3):205-42.

20. Jenkins NW, Parrish JM, Hrynewycz NM, Brundage TS, Singh K. Longitudinal evaluation of patient-reported outcomes measurement information system for back and leg pain in minimally invasive transforaminal lumbar interbody fusion. *Neurospine* 2020;17(4):862-70.

21. Landis JR, Koch GG. The measurement of observer agreement for categorical data. *Biometrics* 1977;33(1):159-74.

22. Pfirrmann CW, Metzdorf A, Zanetti M, Hodler J, Boos N. Magnetic resonance classification of lumbar intervertebral disc degeneration. *Spine (Phila Pa 1976)* 2001;26(17):1873-8.

23. Imagama S, Kawakami N, Matsubara Y, Kanemura T, Tsuji T, Ohara T. Preventive effect of artificial ligamentous stabilization on the upper adjacent segment impairment following posterior lumbar interbody fusion. *Spine (Phila Pa 1976)* 2009;34(25):2775-81.

24. Okuda S, Nagamoto Y, Matsumoto T, Sugiuara T, Takahashi Y, Iwasaki M. Adjacent segment disease after single-segment PLIF: the risk factor for degeneration and its impact on clinical outcomes. *Eur Spine J* 2011;20(11):1946-50.
26. Kim JS, Jung B, Lee SH. Instrumented minimally invasive spinal-transforaminal lumbar interbody fusion (MIS-TLIF): minimum 5-year follow-up with clinical and radiologic outcomes. *Clin Spine Surg* 2018;31(6):E302-9. [PUBMED] [CROSSREF]

27. Park Y, Ha JW, Lee YT, Sung NY. Minimally invasive transforaminal lumbar interbody fusion for spondylolisthesis and degenerative spondylosis: 5-year results. *Clin Orthop Relat Res* 2014;472(6):1813-23. [PUBMED] [CROSSREF]

28. Bin Abd Razak HR, Dhoke P, Tay KS, Yeo W, Yue WM. Single-level minimally invasive transforaminal lumbar interbody fusion provides sustained improvements in clinical and radiological outcomes up to 5 years postoperatively in patients with neurogenic symptoms secondary to spondylolisthesis. *Asian Spine J* 2017;11(2):204-42. [PUBMED] [CROSSREF]

29. Min JH, Jang JS, Lee SH. Comparison of anterior- and posterior-approach instrumented lumbar interbody fusion for spondylolisthesis. *J Neurosurg Spine* 2007;7(1):21-6. [PUBMED] [CROSSREF]

30. Nakai S, Yoshizawa H, Kobayashi S. Long-term follow-up study of posterior lumbar interbody fusion. *J Spinal Disord* 1999;12(4):293-9. [PUBMED] [CROSSREF]

31. Ha DH, Kim TK, Oh SK, Cho HG, Kim KR, Shim DM. Shim D-MiCOS. Results of decompression alone in patients with lumbar spinal stenosis and degenerative spondylolisthesis: a minimum 5-year follow-up. *Clin Orthop Surg* 2020;12(2):187-93. [PUBMED] [CROSSREF]

32. Lee JC, Kim Y, Soh JW, Shin BI. Risk factors of adjacent segment disease requiring surgery after lumbar spinal fusion: comparison of posterior lumbar interbody fusion and posterolateral fusion. *Spine (Phila Pa 1976)* 2014;39(5):E339-45. [PUBMED] [CROSSREF]

33. Min JH, Jang JS, Lee SH. Comparison of anterior- and posterior-approach instrumented lumbar interbody fusion for spondylolisthesis. *J Neurosurg Spine* 2007;7(1):21-6. [PUBMED] [CROSSREF]

34. Nakashima H, Kawakami N, Tsuji T, Ohara T, Suzuki Y, Saito T, et al. Adjacent segment disease after posterior lumbar interbody fusion: based on cases with a minimum of 10 years of follow-up. *Spine (Phila Pa 1976)* 2015;40(14):E831-41. [PUBMED] [CROSSREF]

35. Maragkos GA, Atesok K, Papavassiliou E. Prognostic factors for adjacent segment disease after L4-L5 lumbar fusion. *Neurosurgery* 2020;86(6):835-42. [PUBMED] [CROSSREF]

36. Ghiselli G, Wang JC, Hsu WK, Dawson EG. L5-S1 segment survivorship and clinical outcome analysis after L4-L5 isolated fusion. *Spine (Phila Pa 1976)* 2003;28(12):1275-80. [PUBMED] [CROSSREF]

37. Kim YE, Goel VK, Weinstein JN, Lim TH. Effect of disc degeneration at one level on the adjacent level in axial mode. *Spine (Phila Pa 1976)* 1991;16(3):331-5. [PUBMED] [CROSSREF]

38. Okuda S, Iwasaki M, Miyauchi A, Aono H, Morita M, Yamamoto T. Risk factors for adjacent segment degeneration after PLIF. *Spine (Phila Pa 1976)* 2004;29(14):1535-40. [PUBMED] [CROSSREF]

39. Rouben D, Casnellie M, Ferguson M. Long-term durability of minimal invasive posterior transforaminal lumbar interbody fusion: a clinical and radiographic follow-up. *J Spinal Disord Tech* 2011;24(5):288-96. [PUBMED] [CROSSREF]

40. Solberg TK, Sætlev A, Sjaavik K, Nygaard ØP, Ingebrigtsen T. Would loss to follow-up bias the outcome evaluation of patients operated for degenerative disorders of the lumbar spine? *Acta Orthop* 2011;82(1):56-63. [PUBMED] [CROSSREF]