Transforaminal Lumbar Interbody Fusion (TLIF) Technique for the Treatment of Degenerative Lumbar Disease

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Abstract

Transforaminal lumbar interbody fusion (TLIF) is a relatively new technique of lumbar arthrodesis via posterior transforaminal approach to the disc, indicated mainly in cases of degenerative disc disease, low grade spondylolisthesis and reoperation for disc herniation, especially when there is indication for interbody fusion and posterior decompression. The aim of the study is to assess and evaluate the outcome of posterior instrumented transforaminal lumbar interbody fusion as a surgical treatment modality for symptomatic lumbar degenerative disease. This was a clinical trial study, including 20 cases of symptomatic degenerative lumbar disease who have failed medical treatment which includes one or more of the following: more than 40-year-old, having radiological evidence of lumbar degenerative disease whether stable or unstable. The duration of the study ranged from 6-12 months. Oswestry Disability Index (ODI) at baseline was ranged between 42 – 80 with a mean value of 61.65±11.380 and it was decreased significantly after 1 year of surgery with a mean value of 35.35±11.417. VAS score at baseline was ranged between 4 – 9 with a mean value of 6.80±1.609 and it was decreased significantly after 1 year of surgery with a mean value of 0.75±0.910. TLIF is a technique which offers a simple, safe and effective treatment for degenerative lumbar spine disorders with great improvement of quality of life in cases with surgery satisfaction.

Keywords: Transforaminal Lumbar Interbody Fusion, Lumbar Disc Disease, Oswestry Disability Index, Degenerative Lumbar Spine.

1. Introduction

Interbody fusion techniques have been developed to preserve the load-bearing capacity of the spine, restore sagittal plane alignment and proper disc height, all of which enhance the potential for fusion. The unilateral transforaminal approach for segmental lumbar interbody fusion was first described in the 1980s by [1] and it was popularized in the late 1990s by Harms and Jezerszky [2].

Transforaminal lumbar interbody fusion (TLIF) is an alternative technique, which avoids the anterior approach and the approach through the spinal canal. Theoretically, it prevents typical complications, such as those seen in anterior and posterior lumbar interbody fusion [3]. This approach offers the advantage that it can easily be performed unilaterally. This results in less destruction of the posterior elements and less gross destabilization of the spine, which will maximize fusion stability. Furthermore, it allows better access to the neuroforamen and reduces the need to manipulate spinal nerve roots. Thus, nerve injuries that may occur during retraction may be avoided.

For those with lumbar stenosis but without spondylolisthesis (deformity), the surgical management has traditionally involved posterior decompressive procedures, including laminectomy or laminotomy, and judicious use of partial medial facetectomies and foraminotomies, with or without discectomy. In patients with evidence of spinal instability, however, in situ posterior lumbar fusion is recommended as a treatment option in addition to decompression in the setting of lumbar stenosis [4].

Secondary indications include recurrent lumbar disc herniation, where extensive bony removal is necessary for exposure of the disc fragments, lateral or massive disc herniations, failed previous lumbar fusions by other techniques, and discogenic low back pain [5].

Although most cases of low back pain are transient and relieved by comfort measures along with temporary activity modification, conservative management remains ineffective in approximately 5% of cases that go on to become chronic and disabling [6], resulting in a need for more aggressive treatment.

Lumbar spinal fusion may be used as a potential adjunct, however, in patients with a herniated disc in whom there is evidence of preoperative spinal deformity. Because lumbar deformity, instability, or even chronic low back pain may occur as a result of a reoperative lumbar discectomy, fusion is often considered in the setting of repeated lumbar disc herniations [4].

2. Subjects and Methods

The study had been conducted at Benha university hospital, orthopedics department and Royal National Orthopaedic Hospital Stanmore London.

2.2 Study Design

This is an intervention type of epidemiologic studies termed retrospective study with randomized controlled clinical trial that had been conducted a long twelve months.

2.3 Target Population

Symptomatic degenerative lumbar disease patients who have failed medical treatment.

2.4 Study Population

Symptomatic degenerative lumbar disease patients who have failed medical treatment attending Benha University hospital.

2.5 Inclusion Criteria

This study had been included 20 cases of symptomatic degenerative lumbar disease who have failed medical treatment which includes one or more of the following: more than 40 year old and having radiological evidence of...
Transforaminal Lumbar Interbody Fusion (TLIF) Technique for the Treatment of lumbar degenerative disease whether stable or unstable.

2.6 Exclusion criteria
Any cases with the following criteria will be excluded: Incomplete radiological documentation, inaccurate radiological documentation before or after the TLIF and Anticipated poor cooperation of the patient.

2.7 Sampling technique
Random allocation of forty above 40 years old, symptomatic degenerative lumbar disease patients, who have failed medical treatment attending benha university hospital and Royal national orthopaedic hospital. Twenty of them meet the inclusion criteria and are recruited for the study group, the rest will be the control group.

Sample size: Case group: Twenty symptomatic degenerative lumbar disease patients who have failed medical treatment and give consent to be subjected to surgical intervention aged above 40 years old.

2.8 Control group
Comparable number of symptomatic degenerative lumbar disease patients who had failed medical treatment and don't give consent to be subjected to surgical intervention aged above 40 years old.

STEPS: 0 /58
Step 1-Pedicle screws insertion

2.9 Methods of diagnosis
All patients were evaluated clinically by history and physical examination. Special attention is directed towards associated neurology, previous spine procedures, gait disturbance, and any change in body habitus and posture. All the patients had standing radiographs of the lumbar spine (Anteroposterior and lateral views). Taking care of the radiological signs of frank instability (lithesis more than 3.5 mm or cobbs angle more than 10 degrees).

Assessment and outcome evaluation had been included
Radiological evaluation, Clinical outcomes are assessed by the patients using the Oswestry Disability Index (ODI) and visual analog scale (VAS) questionnaires, Patient satisfaction and clinical improvement and Presence of complications.

2.10 Surgical technique
Antibiotics were given intravenously at induction of anesthesia and for 3 day postoperatively. Then oral antibiotics for 10 post operatively.

2.11 Operative details
- Date of operation - Operative time - Theatre - Blood loss - Anesthesia - Approach - Soft tissue and muscle condition

Fig (1) Identify pedicle insertion points. The optimal insertion point is at the intersection of the transverse process and pars interarticularis [7].

Fig (2) The inferior lamina of L5 can be removed by a kerrison rongeur illustrated by the dotted line [7].

Fig (3) Resect the inferior articular process of L5 with a straight osteotome or a Kerrison [7].
Step 3: Annulotomy and Initial Disc Dissection

Fig (4) Dissector or nerve root retractoris used to ensure the protection of these structures at every step of the procedure [7].

Step 4: Initial Distraction and Preparation of Disc Space

Fig (5) Once distraction is obtained, the opening of the disc space can be maintained with either a temporary rod or the use of a laminar spreader between the spinous processes [7].

Step 5: Final Disc Preparation and Endplate Cleaning

Fig (6) The final discetomy is performed using a combination of curettes, osteotomes, rongeurs and shavers [7].

Step 6: Decortication and Placement of Bone Graft

Fig [7] In order to achieve a solid interbody fusion, the disc space should be filled with as much bone graft as possible. Fill the anterior third and contra-lateral side of the disc space with bone graft using a variety of straight and curved bone tamps from the disc preparation set [7].
Step 7: Cage Trialing

Fig (8) A cage trial should be used prior to insertion of the implant to evaluate potential cage placement and determine the optimal implant fit [7].

STEP 8: Cage Insertion – Concorde Bullet

Fig (9) Align threaded hole of cage with threaded tip. Tighten the knob clockwise until cage is secure. Take care not to cross thread or overtighten the inserter [7].

STEP 9: Implant orientation for lordotic CONCORDE Bullet
STEP 10: Final Compression
STEP 11: Verification of final cage placement – concorde Bullet
An X-ray should be taken to verify final cage placement.

Fig (34) The appearance of three tantalum beads will identify the position of the CONCORDE Bullet cage in the sagittal, coronal and axial planes [7].
Postoperative care

The patients are admitted to the hospital. The patients receive intravenous antibiotics, pain medication as required.

The patient is typically mobilized out of bed the day after surgery.

Follow up

Patients were asked to return to hospital for follow-up at 4 weeks, 3 months, 6 months, 12 months, and thereafter once a year after operation.

Administrative design

- Approvals: - An informed verbal consent from all participants was taken and confidentiality of information was assured. - An official written administrative permission letter was obtained from dean of faculty of medicine, Benha university hospital. The title and objectives of the study were explained to them to ensure their cooperation.

- Ethical committee: Permission from the faculty of medicine ethical committee was also obtained and approval from institutional review board was taken.

3. Results

Table (1) shows TLF level of the studied group show that degenerated segment was L4/L5 in 20(100%), degenerated segment was L5/S1 in 5(25%) and 3(15%) had in multi levels.

Table (2) show operation data results. Surgical time was ranged between 1.5 – 3.5 hours with a mean value of 2.60±0.620 hours. Blood loss was ranged between 160 – 345 ml with a mean value of 248.75±62.993 ml. Hospital stay was ranged between 4 – 7 with a mean value of 5.95±1.146.

Table (3) show assessment tools results. Oswestry Disability Index (ODI) at baseline was ranged between 42 – 80 with a mean value of 61.65±11.380 and it was decreased significantly after 1 year of surgery with a mean value of 35.35±11.417. VAS score at baseline was ranged between 4 – 9 with a mean value of 6.80±1.609 and it was decreased significantly after 1 year of surgery with a mean value of 0.75±0.910.

Table (4) shows Patient satisfaction of the studied group show that 17(85%) had good rate of satisfaction and 3(15%) had poor satisfaction.

Table (5) shows complications of the studied group show that 3(15%) had Infection, 2(10%) had Persistent back pain, 1(5%) had Pseudarthrosis and 1(55) had Neurological deficit.

Table (6) shows Pain improvement of the studied group show that 18(90%) had pain improving and 2(10%) had no improving.
A clinical trial study was held, including 20 cases of symptomatic degenerative lumbar disease who have failed medical treatment which includes one or more of the following: more than 40-year-old, having radiological evidence of lumbar degenerative disease whether stable or unstable. The duration of the study ranged from 6-12 months.

The present study shows that TLF level of the studied group show that degenerated segment was L4/L5 in 20(100%) and degenerated segment was L5/S1 in 5(25%).

[8] demonstrated that clinical presentations in their study were mechanical back pain in 100% of cases and leg pain in 90% of cases. Female to male ratio was 1:4.

Lumbar spinal fusion was introduced approximately 70 years ago and has evolved as a treatment option for symptomatic degenerative lumbar disease.

As regard Pain improvement of the studied group show that 17(85%) had pain improving and 2(10%) had no improving.

Our results are supported by study of [9] as they concluded that the TLIF showed to be a good alternative to PLIF with relatively less risk of complications, less operating time and hospitalization, as well as significant reduction in blood loss during operation. TLIF approach lessens the potential for nerve root injury, therefore resolving probably the most important limitation of the PLIF procedure. For this reason, and the case specific advantages of TLIF over a combined anterior and posterior single-level fusion, they favored TLIF over PLIF as the choice surgical procedure for the posterior operative management of symptomatic degenerative lumbar spine disorders.

Furthermore, [10] reported that Statistically significant differences in favor of minimally invasive treatment were immediate postoperative pain (VAS at third day after operation 4.5 vs. 7.2, P < 0.001), short-term postoperative pain and function (VAS at 30th day after operation 3.2 vs. 5.6, P < 0.001 and ODI 18% vs. 32%, P < 0.001, respectively), hospital stay (4.1 vs. 7.4 d, respectively, P = 0.015), and total blood losses (230 vs. 620 mL, respectively, P < 0.0001). In contrast, there are findings in their study that divert from other evidence in the literature. Park and Ha , [11] as well as [12] report significantly higher surgical time with minimally invasive procedure.

As regard Pain improvement of the studied group show that 18(90%) had pain improving and 2(10%) had no improving.

Our results are supported by study of [13] as they mentioned that lumbar pain improved in 83.5% of patients compared to 90% lumbar pain improvement in [8] study.

Z. Audat et al.[14] reported that pain symptoms relieved in 70% of 81 patients, and good outcomes were reported in 80% of the patients. Show that 17(85%) had good rate of satisfaction and 3(15%) had poor satisfaction. Pain improvement of the studied group show that 18(90%) had pain improving and 2(10%) had no improving. As regard return to work of the studied group show that 1(5%)

Transforaminal Lumbar Interbody Fusion (TLIF) Technique for the Treatment of Degenerative Disc Disease (DDD), internal disc disruption, laminectomy alone when there is evidence of preoperative lumbar spinal deformity or instability that could worsen after procedure.

This is why this study was selected to be conducted to assess and evaluate the outcome of posterior instrumented transforaminal lumbar interbody fusion as a surgical treatment modality for symptomatic lumbar degenerative disease. A clinical trial study was held, including 20 cases of symptomatic degenerative lumbar disease who have failed medical treatment which includes one or more of the following: more than 40-year-old, having radiological evidence of lumbar degenerative disease whether stable or unstable. The duration of the study ranged from 6-12 months.

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didn’t return to work, 15(75%) return to their work and 4(20%) return to work but with light work.

Our results are supported by study of [15] as they reported in their study underwent TLIF surgery that fusion rate radiologically was 95% of cases and good to excellent clinical outcomes was achieved in 88% of cases. In the study of [8], fusion rate was 90% and improvement in clinical symptoms was 90% of cases.

C.L. Goldstein et al. [16] found a dural injury rate of 5.4%, graft malposition of 4.4%, screw mal-position of 2.6%, neurologic deficit and nerve injury of 3.8%, reoperation ratio of 3.3%, and reoperation for graft mal-position of 1.8% for PLIF and TLIF procedures.

K. Asil et al. [17] reported that in his study overall complication rate was 23.9%, dural injury rate was 9.9%, graft mal-position rate was 2.82%, and the screw mal-position rate was 4.23%.

Regarding Z. Audat et al. [14] there was a significant decrease in the ODI (Oswestry Disability Index) scores over time (p < 0.005) but no significant difference among the groups at different follow-up times. Radiographic fusion rates for Groups I, II and III were 88%, 88.9% and 91.9%, respectively.

F. Shunwu et al. [12] reported that the minimally invasive TLIF group was found to have reduced blood loss, fewer transfusions, less postoperative back pain, lower serum creatine kinase on the third postoperative day, a shorter time to ambulation, and a brief hospital stay. The Oswestry Disability Index and Visual Analog Scale scores were significantly lower in the minimally invasive group during follow-up. However, the open group had a shorter operative duration. The complications in the 2 groups were similar, but 2 cases of screw malposition occurred in the minimally invasive group.

5. Conclusion

TLIF is a technique which offers a simple, safe and effective treatment for degenerative lumbar spine disorders with great improvement of life quality of cases with surgery satisfaction.

References


