



2 **Does a high BMI affect the outcome of minimally invasive TLIF?**
3 **A retrospective study of 207 patients**

4 Ayush Sharma¹ · Akash Shakya¹ · Vijay Singh¹ · Nilesh Mangale¹ · Ghanshyam Kakadiya² · Ajay Jaiswal¹ ·
5 Nandan Marathe³

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8 **Abstract**

9 **Purpose** We investigated whether a high Body Mass Index (BMI) affects the outcomes following Minimally Invasive TLIF
10 (MI-TLIF) for degenerative lumbar pathologies.

11 **Methods** A retrospective study was undertaken to include patients operated between January 2016 and January 2020 with at
12 least one-year follow-up. Various preoperative and demographic parameters were recorded and the patients were classified
13 into normal, overweight and obese based on the BMI. The operative and outcome measures used for assessment were surgi-
14 cal time, blood loss, number of levels operated upon, skin incision length, day of independent mobilisation, total hospital
15 stay including ICU stay, return to work and Visual Analogue Score (VAS) for back pain (VAS-BP) and leg pain (VAS-LP)
16 and Oswestry Disability Index (ODI). Attainment of Minimal Clinically Important Difference (MCID) for the scores was
17 calculated. Multivariate analyses were done to assess the effect of BMI on different parameters.

18 **Results** Blood loss and postoperative ICU stay were found to be higher in the obese patients. However, the other variables
19 were comparable. VAS-BP, VAS-LP and ODI scores were significantly improved in all the patients with no inter-group
20 variability. The MCID attainment was also similar. The satisfaction rating at 1-year and willingness for surgery again for
21 similar disease was also similar. The overall complication rate was 14.9% and was comparable among the groups. Multivari-
22 ate analyses revealed no significant association between BMI and various parameters.

23 **Conclusion** In patients treated by MI-TLIF for degenerative lumbar spine pathology, BMI is not a factor that negatively
24 affects the functional and clinical outcomes.

25 **Keywords** MI-TLIF · Obesity · ODI · MCID · VAS · Outcomes

26 **Introduction**

27 Obesity is an epidemic of the modern ages and has a sig-
28 nificantly high prevalence in the developed world and along
29 with being overweight, has spread to involve approximately
30 a third of the world population [1]. It is predicted that if
31 current trends are to be extrapolated, 38% of the adult popu-
32 lation of the world will be overweight and a further 20%

obese by the year 2030 [2]. Obesity is defined as a body
mass index (BMI) ≥ 30 kg/m² and overweight is defined
as a BMI ≥ 25 kg/m² [1]. This has obvious medical effects
on the body including non-communicable diseases. It also
has particularly negative repercussions on the health of the
spine, especially the lumbar spine and has been well docu-
mented. [3, 4]. The increased incidences of low backaches,
disc degenerations and other similar degenerative conditions
requiring treatment provide the proof needed and also give
the necessary warning as to what might present to us in the
future [5].

The surgery for lumbar degenerative disc disease (LDDD)
commonly involves procedures to cause interbody fusion.
Several approaches and procedures have been described
to attain this with Transforaminal interbody fusion (TLIF)
among one of them. Although the open surgeries were rela-
tively successful in achieving the fusion and improving the

A1 ✉ Akash Shakya
A2 akashshakya.gmc@gmail.com

A3 ¹ Department of Orthopaedics and Spine Surgery, Dr BAM
A4 Hospital, Mumbai, India

A5 ² Department of Orthopaedics and Spine Surgery, BYL Nair
A6 Hospital, Mumbai, India

A7 ³ Department of Orthopaedics and Spine Surgery, KEM
A8 Hospital, Mumbai, India

50 functional outcomes, they carried with them many disadvantages like a longer incision and surgery time and more blood
51 loss. There was also a delayed mobilisation, longer hospital
52 stay and delayed return to work observed. [6] Hence, such
53 surgeries via the minimally invasive route e.g. Minimally
54 Invasive TLIF (MI-TLIF) have attained a lot of traction due
55 to the benefits like a lesser postoperative pain and early reha-
56 bilitation [6]. These are thus comparable to, if not better,
57 than the traditional open approach. In addition, the compli-
58 cations seen in patients with obesity can have an important
59 part to play in the process of surgical decision-making [7].

60 The presence of obesity in a patient requiring surgery
61 for LDDD brings with it, its own set of challenges. Complications thus arising have been studied extensively and
62 described in detail. Functional outcomes after surgeries in
63 obese and overweight patients following open surgeries have
64 shown mixed results. The additional challenges mainly relating to the extensive dissection required to achieve a proper
65 exposure were supposed to be responsible. While a few have
66 reported poorer results, others have observed no similar differences. [8–10] Minimally invasive surgeries have been,
67 however, studied scarcely with respect to this category of
68 patients [11–13]. So, the authors felt the need to evaluate
69 and compare the results of MI-TLIF in different BMI categories and to infer if there was any significant difference
70 in the results. Hence, this study aims to investigate whether
71 obesity and overweight affect the functional outcomes following MI-TLIF in patients undergoing surgery for LDDD
72 compared to those with normal BMI.
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Methodology

79 The retrospective study was conducted at a tertiary care hos-
80 pital in Western India. The patients' data were recorded in
81 the case record forms from the hospital data. The patients
82 operated with MI-TLIF for LDDD from January 2016 to
83 January 2020 were enrolled in the study as per the inclu-
84 sion and exclusion criteria Table 1 All the relevant demo-
85 graphic data and preoperative variables were collected at the
86 time of admission. The patients were classified into three
87 groups based on their BMI i.e. normal ($\text{BMI} < 25 \text{ kg/m}^2$),
88 Overweight ($25\text{--}29.9 \text{ kg/m}^2$) and Obese ($\geq 30 \text{ kg/m}^2$). The
89 intraoperative data were collected from the surgery records.
90 The postoperative data were collected on postoperative day
91 1, 1 month, 3 months, 6 months and 1 year and then every
92 year as a routine for all the patients.
93

Surgical technique (Fig. 1 and Fig. 2)

94 All the surgeries were performed by the same team of sur-
95 geons. Under general anaesthesia, the patient was positioned
96 prone and a 22 mm tubular retractor system docked on the
97 facet joint of the symptomatic side under radiographic guid-
98 ance as per the Wiltse paraspinous approach. Removal of the
99 inferior facet followed by the superior facet was done using
100 an osteotome or ultrasonic scalpel. Flavectomy was done
101 and the neural structures were identified. Discectomy was
102

Table 1 Inclusion and exclusion criteria for the patient selection

Inclusion criteria	Exclusion criteria
Age ≥ 18 years	Previous surgery of the lumbar spine
Degenerative lumbar pathology	Other pathologies like trauma, tumour, infection
Follow up ≥ 12 months	Patients with spondyloarthropathies
Willingness for participation in the study	Concomitant cervical/dorsal spine pathology



Fig. 1 Procedure of the MI-TLIF by a case example a Preoperative MRI of the lumbosacral spine in the sagittal plane of a male patient with normal BMI having a degenerative disc disease at L4-L5 and L5-S1; b, c: Intraoperative image-intensifier images showing cage implantation at L4-L5 level as seen in the lateral b and anteroposterior c views; d, e: Completion of the procedure with cage implan-

tation at L5-S1 and pedicle screw and rod placement as seen in the lateral d and anteroposterior e views; f and g: Radiographs of the lumbosacral spine in the anteroposterior f and lateral g views at one-year follow up of the same patient showing a good fusion at both the levels

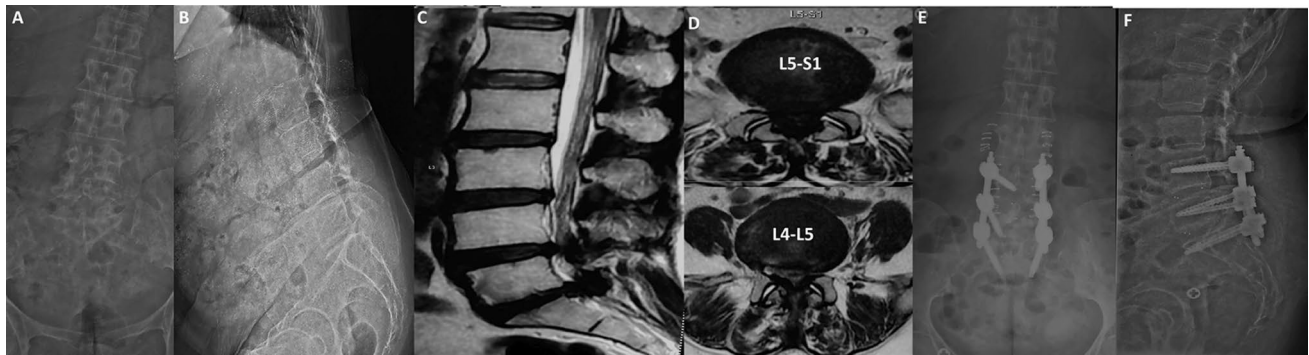


Fig. 2 Example MI-TLIF in an obese female a, b: Preoperative radiographs of the lumbosacral spine in the anteroposterior a and lateral b views of the patient showing degenerative disease with loss of lumbar lordosis; c, d: Preoperative MRI of the lumbosacral spine showing

severe lumbar canal stenosis at L4-L5 and L5-S1 in the sagittal c and axial planes d; e and f: Postoperative radiographs of the lumbosacral spine in the anteroposterior e and lateral f views showing MI-TLIF at L4-L5 and L5-S1

103 completed and adequate endplate preparation done was with
 104 curettes. The morselised autograft obtained was packed
 105 anteriorly under the anterior longitudinal ligament and in
 106 the PEEK cage which was then implanted. Fig. 1b, c Over-
 107 the-top decompression was done in every case and adequate
 108 decompression was ensured. Facetectomy on the opposite
 109 side was not done in any case. Percutaneous titanium pedicle
 110 screw insertion and rod placement after appropriate compression
 111 completed the procedure Fig. 1d, e.

112 The operative and outcome measures used for assessment
 113 were the surgical time, blood loss, number of levels operated
 114 upon, skin incision length, day of independent mobilisation,
 115 total hospital stay including ICU stay if any, return
 116 to work and Visual Analogue Score (VAS) for back pain
 117 (VAS-BP) and leg pain (VAS-LP) and Oswestry Disability
 118 Index (ODI). Whether the improvement obtained in the ODI,
 119 VAS-BP and VAS-LP scores was meaningful was ascertained
 120 by the attainment of the Minimal Clinically Important
 121 Difference (MCID) for each variable. So, the final outcome
 122 was dichotomised based on whether the patient achieved the
 123 MCID threshold or not. This was taken as 14.9 for the ODI
 124 score and 2.1 and 2.8 for VAS-BP and VAS-LP, respectively
 125 [14].

126 The fitness for returning to work was assessed by an
 127 independent Occupational Therapist who was blinded to the
 128 intervention. The patients were also asked to rate their satisfaction
 129 at the end of 1 year by choosing a number between 0
 130 and 10 with 0 representing “completely unsatisfied” and 10
 131 representing “completely satisfied”. They were also asked at
 132 the end of 1 year whether they would like to have the procedure
 133 for the same pathology in the future and the responses
 134 were recorded as “yes”, “no” and “not sure”. The fusion
 135 rates were assessed at the end of 1 year based on the findings
 136 on the computed tomographic (CT) imaging.

137 The differences between study groups were tested using
 138 One-way analysis of variance (ANOVA) with equal variance

assumptions for continuous variables and Chi-Squared tests
 with Yates’ correction for continuity. To determine the
 effects of BMI on surgery time, a multivariate regression
 was performed. Similar analyses were also performed to
 evaluate the effect of BMI on blood loss, hospital stay, incision
 length, operative time, return to work and changes in the
 VAS and ODI scores.

Results

146 A total of 207 consecutive patients were enrolled in the
 147 study of which 33 were obese, 53 overweight and the rest
 148 121 patients had a BMI < 25. There was no significant difference
 149 between the three groups in terms of gender, age, pre-
 150 operative symptom duration, diagnosis and baseline clinical
 151 scores. The prevalence of diabetes mellitus and dyslipidaemia
 152 was significantly higher in the obesity group although
 153 no difference was seen in the incidence of other pre-existing
 154 disorders Table 2.

155 In terms of the clinical and operative variables, the number
 156 of levels operated upon, total hospital stay, operative time
 157 and the cumulative size of the skin incisions were found to
 158 be comparable ($p > 0.05$). Table 3 However, blood loss was
 159 significantly more in the obese patients ($p = 0.006$) as was
 160 the ICU stay postoperatively ($p = 0.03$). The patients were
 161 mobilised at 3.6 ± 1.1 days postoperatively. Though patients
 162 with normal BMI were mobilised earlier (3.5 ± 1.1 days) as
 163 compared to those in the overweight (3.7 ± 1.2 days) and
 164 obese (4.0 ± 1.0 days) groups, no significant difference was
 165 observed among the different groups ($p = 0.07$). Return to
 166 work was similarly comparable with the patients joining at
 167 an average of 6.8 ± 1.1 weeks after the surgery ($p = 0.32$).
 168

169 Functional assessment was done using ODI scores and
 170 VAS-BP and VAS-LP scores. ODI scores were significantly
 171 better at 6 months and 1-year postoperatively as compared

Table 2 Demographic variables of the patients

Variable	Normal (n=121)	Overweight (n=53)	Obese (n=33)	P Value
Age (years)	51.9	53.9	56.6	0.09
Mean	11.8	10.4	9.3	
Standard Deviation				
Gender	77	30	21	0.66
Male	44	23	12	
Female				
BMI (kg/m ²)	22.3	27.7	33.3	< 0.001
Mean	1.6	1.5	1.6	
Standard Deviation				
Duration of Symptoms (months)	7.4	7.8	8.2	0.12
Mean	2.1	1.9	2.4	
Standard Deviation				
Diagnosis	40	22	15	0.84
Lumbar Canal Stenosis	13	4	3	
Degenerative Spondylolisthesis	43	19	12	
Degenerative Disc Disease	17	6	2	
Spinal Instability	8	2	1	
Degenerative Scoliosis				
Follow up (months)	15.8	16.2	14.9	0.30
Mean	3.9	4.1	2.9	
Standard Deviation				
Comorbidities	20	13	10	0.10
Hypertension	14	14	5	< 0.05
Diabetes Mellitus	4	2	4	0.15
Cardiac Disease	5	4	2	0.50
COPD	7	2	1	0.50
Thyroid Disorder	21	18	19	< 0.0001
Dyslipidaemia	13	10	4	0.37
Smoking				

A *p* value < 0.05 is statistically significant

172 to the pre-operative scores in all the groups and the scores
173 of all the three groups were comparable. ($p=0.84$) Table 4.
174 Similarly, VAS-BP and VAS-LP were significantly improved
175 at the end of 1 year as compared to the preoperative scores
176 ($p<0.001$) with no significant inter-group difference at
177 any point of assessment Table 5. On being asked at 1 year,
178 whether in hindsight, the patients would like to have the
179 same treatment for their presenting complaints, the results
180 were comparable ($p=0.60$). Table 3 Similarly, the average
181 satisfaction rating after the surgery was 8.8 ± 1.1 with no
182 significant difference among the groups ($p=0.42$) Table 3.

183 A multivariate linear regression analysis was done to
184 determine the effect of BMI on the improvement in ODI
185 and VAS scores at 1 year compared to the preoperative lev-
186 els controlling for age, sex, the number of levels fused and
187 comorbidities. It was found that BMI was not associated
188 with the changes in either the ODI score ($p=0.84$) or the
189 VAS scores ($p=0.33$). Similar analyses to evaluate the effect

of BMI on hospital stay ($p=0.48$), blood loss ($p=0.72$),
operative time ($p=0.73$) and size of the skin incisions
($p=0.80$) were found to be non-significant.

The complications encountered were divided into two
groups i.e. early and late depending on the occurrence
before and after six months of the surgery. Early complica-
tions were further subclassified into dural tears and non-
dural tears. In the group of early complications, 20 cases of
accidental durotomies and 8 cases of non-dural tear com-
plications were witnessed Table 3. Two cases of dural tears
required re-exploration with closure of the defect within a
week of the surgery. With respect to the non-dural tear com-
plications, there were two episodes of superficial infections
which resolved on conservative therapy. There was one epi-
sode of urinary tract infection which was treated with anti-
biotics. One patient developed postoperative pneumonia which
was managed in the ICU for one week. Two patients experi-
enced worsening of symptoms within a month of surgery of

Table 3 Operative and outcome variables of the patients

Variable	Normal BMI (N = 121)		Overweight (N = 53)		Obese (N = 33)		p Value
	Mean	Standard Deviation	Mean	Standard Deviation	Mean	Standard Deviation	
Duration of Surgery (minutes)	99.4	19.4	96.7	20.8	103.0	18.7	0.35
Blood Loss (millilitres)	95.5	27.0	91.0	29.5	111.1	32.1	0.006
Incision Length (mm)	36.1	2.5	35.7	2.7	36.3	2.7	0.52
Hospital Stay (days)	8.9	3.2	9.7	3.5	10.2	3.4	0.09
ICU Stay	0.8	1.1	0.9	1.3	1.4	1.1	0.03
Day of Independent Mobilisation	3.5	1.1	3.7	1.2	4.0	1.0	0.07
Return to Work (weeks)	6.7	1.1	6.9	1.2	7.0	1.3	0.32
Number of Levels	1	95	41		26		0.98
	2	24	11		6		
	3	2	1		1		
Complications	DT	10	6		4		0.72
	NDT	5	3		3		0.53
	Total	15	9		7		0.40
Patients requiring re-surgery within one year	4		1		1		0.88
Willingness for surgery again	Yes	101	39		25		0.60
	No	9	6		3		
	Not Sure	11	8		5		
Satisfaction Rating	8.9	1.0	8.7	1.3	8.7	1.1	0.42
MCID Attainment	VAS-BP	103 (85.1%)	46 (86.8%)		29 (87.8%)		0.90
	VAS-LP	105 (86.8%)	45 (84.5%)		28 (84.5%)		0.93
	ODI	110 (90.9%)	43 (81.1%)		28 (84.5%)		0.18
Fusion Attained	115 (95.0%)		49 (92.4%)		30 (90.9%)		0.62

ICU Intensive Care Unit, DT Dural tear, NDT Non-Dural Tear, MCID Minimal Clinically Important Difference, VAS Visual Analogue Score, BP Back pain, LP Leg pain, ODI Oswestry Disability Index

A p value < 0.05 is statistically significant

Table 4 Comparison of Oswestry Disability Index (ODI) scores at the various time intervals among the three groups

Group	Preoperative	Postoperative Day 1	Postoperative 1 month	Postoperative 3 months	Postoperative 6 months	Postoperative 1 year	Significance of the improvement in ODI score at 1 year (P Value)
Normal	32.2	25.5	19.1	16.4	15.0	12.0	< 0.001
	Mean	3.6	2.9	2.4	3.0	1.8	
Overweight	33.4	25.0	19.9	17.0	14.8	11.8	< 0.001
	Mean	3.9	3.4	3.3	2.1	2.0	
Obese	32.2	24.4	19.1	17.2	14.9	12.0	< 0.001
	Mean	4.6	3.0	3.0	2.1	1.8	
P Value	0.15	0.16	0.19	0.19	0.80	0.84	

ODI: Oswestry Disability index

A p value < 0.05 is statistically significant

208 which one required surgery and the other resolved on conservative
209 management within 3 months. Two patients developed a postoperative neurological deficit of which 1 was
210

managed with revision surgery within two weeks and the other improved gradually over 6 months. Regarding the late complications, three patients developed recurrent symptoms

211
212
213

Table 5 Comparison of Visual Analogue Score (VAS) scores at the various time intervals among the three groups

Group	Preoperative		Postoperative day 1		Postoperative 1 month		Preoperative		Postoperative 3 months		Postoperative 6 months		Postoperative 1 year		Significance of improvement in VAS score at 1 year (<i>P</i> Value)
	BP	LP	BP	LP	BP	LP	BP	LP	BP	LP	BP	LP	BP	LP	
Normal Mean	7.5	6.5	6.2	6.0	5.5	5.5	5.3	5.0	4.9	4.3	4.3	4.3	3.6	3.1	<0.001
Standard Deviation	0.8	1.1	1.1	1.3	1.2	1.3	1.3	1.4	1.4	0.9	0.9	1.0	1.0	0.7	<0.001
Overweight Mean	7.4	6.9	6.5	6.1	5.9	5.8	4.9	4.9	4.1	4.5	4.5	3.6	2.9	1.3	<0.001
Standard Deviation	1.0	1.2	0.9	1.0	0.8	1.0	1.1	1.0	1.2	1.3	1.3	1.1	1.1	1.3	<0.001
Obese Mean	7.2	6.6	6.6	6.3	5.7	5.9	5.0	5.3	4.2	4.0	4.0	3.8	3.2	1.0	<0.001
Standard Deviation	1.1	1.2	1.2	1.1	1.0	1.2	0.9	1.2	0.7	1.2	1.2	1.0	1.0	1.0	<0.001
<i>P</i> Value	0.24	0.11	0.96	0.44	0.08	0.14	0.10	0.36	0.56	0.11	0.59	0.29	0.29	0.29	

VAS: Visual Analogue Score, LP: Leg pain, BP: Back pain

A *p* value < 0.05 is statistically significant

214 after 6 months of surgery and were offered revision surgery.
 215 Two consented for the re-surgery and were operated upon.
 216 However, one refused to give consent for the surgery and
 217 hence was managed conservatively.

218 At the end of the one-year follow-up, a total of 6 patients
 219 were re-operated of which four had normal BMI and one
 220 each belonged to the overweight and obese cohorts. The
 221 distribution of the same across the three groups was non-
 222 significant ($p = 0.36$) Table 3. The mean fusion rate at the
 223 end of 1 year was 93.7% and was comparable among the
 224 groups ($p = 0.62$). The overall complication rate in our
 225 study was 14.9%.

226 MCID assessment was done to determine if the interven-
 227 tion resulted in any meaningful improvement in the func-
 228 tional outcomes as measured by ODI, VAS-BP and VAS-LP
 229 scores. The attainment of the MCID thresholds for all three
 230 variables at the end of one year was comparable Table 3.

231 Discussion

232 Obesity presents a whole new range of problems in the sur-
 233 gical management of the patients requiring lumbar interbody
 234 fusions. These can be either due to the medical or the surgi-
 235 cal issues relating to the patient [15]. The surgical technique
 236 also presents numerous challenges including positioning,
 237 excessive retraction during exposure and ease of image guid-
 238 ance. Many authors have presented poor outcomes following
 239 surgery in these patients [8, 16, 17]. In addition, there are
 240 risks for medical complications after the surgery like pul-
 241 monary embolism, myocardial infarction and gastric ulcers
 242 [15]. These observations thus can bias the surgeon's decision
 243 on the need for surgery for the patient.

244 Obesity is a risk factor for various metabolic conditions
 245 as was observed in our study with diabetes mellitus and dys-
 246 lipidaemia significantly more in the obese patients. The sig-
 247 nificantly high postoperative ICU admission in those having
 248 a BMI ≥ 30 kg/m² seen in our study can also be explained by
 249 the requirement of care for the management of these comor-
 250 bidities. However, a majority of the operative variables were
 251 comparable across the three groups except for the intraop-
 252 erative blood loss. The tubular-assisted MI-TLIF renders
 253 the need for a wide and extensive exposure for obtaining a
 254 proper visualisation meaningless. This observation is similar
 255 to those made by Goh et al. [13]. The increased depth that
 256 is needed to be negotiated in an open approach is bypassed
 257 in the MI approach by the direct docking of the tubular sys-
 258 tem on the target area. Also, the fat and other soft tissues
 259 seem to provide additional stability to the entire tubular
 260 system. Despite the comparable surgical duration, incision
 261 length and number of levels fused, the mean blood loss was
 262 significantly higher in obese patients. This could be due to

an increased venous pressure secondary to the raised intra-
 abdominal pressure on positioning the patient prone for the
 surgery.

In our study, the patients were mobilised independently at
 a mean of 3.6 ± 1.1 days postoperatively which was compa-
 rable among the three groups. This demonstrates that a high
 BMI is not an obstacle for early mobilisation, as the mini-
 mally invasive approach led to a lesser tissue destruction and
 hence a lesser postoperative pain even in high BMI patients.
 A similar trend was observed with respect to the hospital
 stay and return to work. Comparable results were seen by
 others [11, 18]. Return to work observed by us was at a
 mean of 6.8 ± 1.1 weeks which was less than that observed
 by Adogwa et al. who reported a median time of 8.5 weeks
 post-surgery [19].

Both the VAS-BP and VAS-LP decreased significantly
 postoperatively in all the patients with no inter-group vari-
 ability at the end of 1 year. A similar improvement was seen
 in the ODI scores Tables 4, 5. Similar results were obtained
 other [11–13]. In contrast, Herold et al. observed signifi-
 cant but inferior improvements in all the three parameters
 in the patients with BMI > 30 kg/m² undergoing microsurgical
 decompression for lumbar spine for spinal stenosis as
 compared to the non-obese patients which were supposed
 to be due to a higher expectation in the obese patients [17].
 Although the improvement in the VAS-BP, VAS-LP and
 ODI scores was significant overall, it does not necessarily
 translate to a significant clinical improvement at an indi-
 vidual level. This is because a higher improvement in a small
 number of patients can offset a smaller improvement in the
 rest of the patients within the group. Hence, the concept of
 MCID was found to be useful which enables the identifica-
 tion of patients with an improvement in the scores necessary
 to produce a significant clinical and functional improvement
 [14]. Accordingly, the patients in each group were classified
 into two subgroups based on the attainment of MCID. The
 MCID attainment was similar in all the groups for VAS-BP,
 VAS-LP and ODI. Thus, similar outcomes can be expected
 irrespective of the BMI status after MI-TLIF for degenera-
 tive lumbar pathology. The similar satisfaction ratings and
 the willingness for surgery again can be attributed to this
 significant finding.

Complications encountered were classified into early and
 late. The overall rate of durotomies was 9.7% which was
 comparable to that seen elsewhere [20]. A comparable inci-
 dence of dural tears was witnessed among the three groups.
 Burks et al. had noted that a higher incidence of incidental
 dural tears was seen in the obese patients which was not
 seen in our study [20]. The overall complication rate in our
 study was 14.9% which was similar to that observed by
 Wong et al. (15.6%) but lower than that observed by Joseph
 et al. (19.2%) [21, 22]. There was also no significant differ-
 ence in the complication rates among the three groups as

316 also reported by others [11]. However, Marques-Lara et al.
317 reported a higher risk of complications like deep venous
318 thrombosis, pulmonary embolism, infections and postop-
319 erative acute renal failure with a BMI > 25 kg/m², with a
320 greater risk with increasing BMI value [23]. The fusion rate
321 observed at the end of one year was 93.7% and was more
322 than the fusion rate of 92.5% seen by Parajón et al. but less
323 than the rate of 94.7% observed by Bevevino et al. [24, 25].

324 There are a few limitations in the study. As this is a single
325 centre study, the results cannot be generalised to the univer-
326 sal population where new confounders may arise. Secondly,
327 a longer minimum follow-up could lead to the emergence
328 of different patterns in results especially with respect to the
329 functional and radiological outcomes. Thirdly, additional
330 functional outcomes measures like SF-36 could be used.

331 Lastly, the study is limited by its retrospective nature.

332 Conclusion

333 In patients treated by MI-TLIF for degenerative lumbar
334 spine pathology, BMI is not a factor that negatively affects
335 the functional and clinical outcomes. Minimally invasive
336 approach can result in better overall outcome in patients with
337 degenerative lumbar disorders with high BMI.

339 **Data availability** The datasets generated during and/or analysed dur-
340 ing the current study are available from the corresponding author on
341 reasonable request.

342 Declaration

343 **Conflict of Interest** The authors declare that they have no conflict of
344 interest.

345 **Ethical Approval** Ethical approval was waived by the local Ethics Com-
346 mittee of the hospital in view of the retrospective nature of the study
347 and all the procedures being performed were part of the routine care.

348 **Consent to participate** All participants gave written consent for par-
349 ticipation in the trial.

350 **Consent to Publish** All participants gave written consent for permis-
351 sion to publish.

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