RESEARCH



A systematic review and meta-analysis of risk factors for reoperation after degenerative lumbar spondylolisthesis surgery

Yuzhou Chen^{1,2}, Yi Zhou³, Junlong Chen⁴, Yiping Luo⁵, Yongtao Wang² and Xiaohong Fan^{2*}

Abstract

Background Considering the high reoperation rate in degenerative lumbar spondylolisthesis (DLS) patients undergoing lumbar surgeries and controversial results on the risk factors for the reoperation, we performed a systematic review and meta-analysis to explore the reoperation rate and risk factors for the reoperation in DLS patients undergoing lumbar surgeries.

Methods Literature search was conducted from inception to October 28, 2022 in Pubmed, Embase, Cochrane Library, and Web of Science. Odds ratio (OR) was used as the effect index for the categorical data, and effect size was expressed as 95% confidence interval (CI). Heterogeneity test was performed for each outcome effect size, and subgroup analysis was performed based on study design, patients, surgery types, follow-up time, and quality of studies to explore the source of heterogeneity. Results of all outcomes were examined by sensitivity analysis. Publication bias was assessed using Begg test, and adjusted using trim-and-fill analysis.

Results A total of 39 cohort studies (27 retrospective cohort studies and 12 prospective cohort studies) were finally included in this systematic review and meta-analysis. The overall results showed a 10% (95%CI: 8%-12%) of reoperation rate in DLS patients undergoing lumbar surgeries. In surgery types subgroup, the reoperation rate was 11% (95%CI: 9%-13%) for decompression, 10% (95%CI: 7%-12%) for fusion, and 9% (95%CI: 5%-13%) for decompression and fusion. An increased risk of reoperation was found in patients with obesity (OR = 1.91, 95%CI: 1.04–3.51), diabetes (OR = 2.01, 95%CI: 1.43–2.82), and smoking (OR = 1.51, 95%CI: 1.23–1.84).

Conclusions We found a 10% of reoperation rate in DLS patients after lumbar surgeries. Obesity, diabetes, and smoking were risk factors for the reoperation.

Keywords Degenerative lumbar spondylolisthesis, Reoperation, Risk factors, Meta-analysis

² Department of Orthopedics, Hospital of Chengdu University

of Traditional Chinese Medicine, No.39 Shi-Er-Qiao Road, Jinniu District, Chengdu 610075, P.R. China

- ³ Department of Traditional Chinese Medicine, The Traditional Chinese
- Medicine Hospital of Wenjiang District, Chengdu 611130, P.R. China

of Wenjiang District, Chengdu 611130, P.R. China



© The Author(s) 2023. **Open Access** This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/. The Creative Commons Public Domain Dedication waiver (http://creativecommons.org/publicdomain/zero/1.0/) applies to the data made available in this article, unless otherwise stated in a credit line to the data.

⁵ Department of Gynecology, Hospital of Chengdu University of Traditional Chinese Medicine, Chengdu 610075, P.R. China

^{*}Correspondence:

Xiaohong Fan

drfxiaohong@163.com

¹ Chengdu University of Traditional Chinese Medicine, Chengdu 610075, P.R. China

⁴ Department of Anorectal, The Traditional Chinese Medicine Hospital

Background

Degenerative lumbar spondylolisthesis (DLS) refers to anterolisthesis of one vertebral body over another vertebral body secondary to osteoarthritic degeneration, leading to spinal canal stenosis [1]. DLS is an aging-related disease, and its incidence is increasing under the background of the global population aging [2]. Each year, 39 million individuals are diagnosed with DLS, accounting for a global prevalence of 0.53% [3]. DLS may be accompanied with low back pain, radiculopathy, or neurogenic claudication [2].

Surgeries have been regarded as the standard treatment modality for intractable cases [4]. The proportion of lumbar surgeries increases by more than two-fold not only because of elevated prevalence of degenerative lumbar spine disease but also because of improved surgical techniques, good outcomes, and increased hospitals and surgeons [5, 6]. However, due to complications (such as fusion failure, persistent pain, and infection), progressive degenerative changes-related diseases, or an unrelated previous surgeries, some patients require reoperation [7]. Despite improvements in surgical skills and techniques, the reoperation rate is still unimproved, with a 10-year reoperation rate of about 15% [7]. Given the high prevalence and chronicity of DLS, understanding the risk factors for reoperation is important [8]. Park et al. have revealed the longitudinal trends in the lumbar reoperation rate, and the reoperation was associated with demographics, comorbidities, primary surgery type, and preoperative spinal pathology [9]. Noh et al. haven found lifestyle-related factors, such as smoking, drinking, and exercise, were associated with the higher rate of reoperation [10]. However, results of studies on the risk factors for reoperation of DLS patients remain controversial. Rabah et al., have reported that diabetes was related to greater risk of reoperation [11], while Khan et al. reported no significant association between diabetes and reoperation [12]. In the study performed by Zhong et al., obesity was found to be associated with a higher incidence of unplanned reoperations [8]. Nevertheless, Kuo et al. found that obesity was not significantly associated with the reoperation [13].

Considering the controversial results, we aimed to perform a systematic review and meta-analysis to evaluate the incidence and risk factors of reoperation in DLS patients for the purpose of improving surgical outcomes and prognosis.

Methods

Literature search strategy

This study was performed based on the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) [14]. Two researchers (YZC and YZ) conducted the literature search from September 28, 2022 to October 28, 2022 in Pubmed, Embase, Cochrane Library, and Web of Science. Consensus was reached by discussion; if consensus cannot be reached by discussion, a third researcher (XHF) was consulted. Search terms were "degenerative spinal diseases" OR "degenerative spondylolisthesis" OR "degenerative lumbar spondylolisthesis" OR "degenerative cervical spondylolisthesis" AND "spinal surgery" OR "fusion" OR "reoperation" OR "repeat surgery" OR "risk factor".

Selection criteria

Studies meeting the following inclusion criteria were selected: (1) population: DLS patients; (2) patients undergoing lumbar surgeries, including decompression surgeries and fusion surgeries; (3) outcome: reoperation rate and risk factors; and (4) studies: cohort studies. The population included DLS patients and DLS patients with lumbar spinal stenosis (LSS). Fusion surgeries included posterolateral lumbar fusion (PLF) and lumbar interbody fusion (LIF). Reoperation was defined as the secondary lumbar surgeries due to progression of lumbar degenerative changes or postoperative instability [8]. Risk factors included body mass index (BMI), sex, age, diabetes, smoking, and more bleeding.

Studies were excluded by meeting one of the following criteria: (1) animal studies; (2) other degenerative spinal diseases including lumbar spine stenosis, degenerative disc disease, and degenerative cervical spondylosis; (3) not English articles; (4) unable to extract data; (5) case reports, conference abstracts, letters, reviews, and meta-analysis.

Data extraction and critical appraisal

The following data were extracted: the first author, year of publication, country, study design, patients, definition of spondylolisthesis, sample size, age, sex, BMI, disease duration, surgery types, follow-up time, number of reoperations, reasons for reoperation, reoperation methods, and risk factors of reoperation. The Newcastle–Ottawa Scale (NOS) was applied to evaluate the quality of cohort studies [15]. This scale consisted of three items: selection, comparability, and outcome. This scale was scored a total of 9 points, and divided into low quality (0–3 points), fair quality (4–6 points), and high quality (7–9 points) [15].

Statistical analysis

All statistical analyses were performed using Stata15.1 software (Stata Corporation, College Station, TX, USA). Rate was used as the effect index in the analysis of reoperation rate. Odds ratio (OR) was used as the effect index for categorical data, and effect size was represented as 95% confidence interval (CI). Heterogeneity test was

performed for each outcome effect size, and results were quantified as I-squared (I²). Random-effect model was used for analysis if $I^2 \ge 50\%$, and fixed-effect model was used if $I^2 < 50\%$. For the high heterogeneity ($I^2 \ge 50\%$), subgroup analysis was conducted based on study design, patients, surgery types, follow-up time, and quality of studies to explore the source of heterogeneity. Sensitivity analysis was carried out for all outcomes. Begg test was used to assess publication bias for the outcome included more than 10 articles. Trim-and-fill analysis was used to adjust the publication bias. P < 0.05 was considered statistically significant.

Results

Identification of studies and characteristics of patients

A total of 7,662 articles were searched from Pubmed (n=1026), Embase (n=1349), Web of Science (n=4441), and Cochrane Library (n=846). After removing the duplicates, 5,251 articles remained. Further, 5,150 articles were excluded due to publishing as reviews or meta-analyses (n=929), conference abstracts (n=310), animal trials (n=22), case reports (n=226), and letters (n=9),

not English articles (n=112), and topic not meeting the requirements (n=3542). In the remaining 101 articles, we further excluded 3 articles unable to extract data, 29 articles reporting other degenerative spinal diseases, and 30 articles with topic not meeting the requirements. Finally, 39 cohort studies were retained in this meta-analysis [7, 8, 10–13, 16–48], with 27 retrospective cohort studies and 12 prospective cohort studies. The flow diagram of our searching was displayed in Fig. 1. In the included studies, 28 studies were assessed as fair quality, and 11 studies were assessed as high quality. Characteristics of the included studies were presented in Table 1.

Overall results and subgroup analysis results of reoperation rate

This meta-analysis showed a 10% (95%CI: 8%-12%) of reoperation rate in DLS patients (Fig. 2). A high heterogeneity was observed in the results ($I^2=99.3\%$). To explore the source of heterogeneity, subgroup analyses were performed. In study design subgroup, 10% of reoperation rate was found in both prospective cohort study (95%CI: 7%-13%) and retrospective cohort study

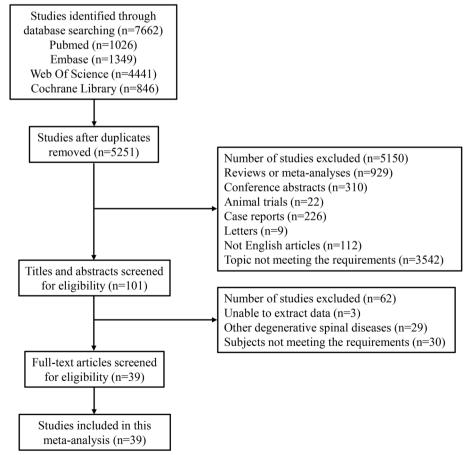


Fig. 1 Flowchart of identifying studies

Table 1 Characteristics of included patients	aracteristics	of included pa	Inerius					
Author	Year	Country	Study design	Patients	Definition of spondylolisthesis	Sample size	Age (years)	Male/female
Salimi	2022	Japan	Retrospective cohort	DLS with LSS	≥3 mm anterior slippage	50	69.4±9.5	23/27
Noh	2022	Korea	Retrospective cohort	DLS with or without LSS	ЧA	DLS with LSS: 3840; DLS without LSS: 255	60.5 ± 9.1	NA
Moayeri	2022	Canada	Retrospective cohort	DLS with LSS	grade 1	140	68.0±10.1	64/76
Liang	2022	China	Retrospective cohort	DLS with LSS	grade 1	104 (PMTD: 53; MIS TLIF: 51)	62.06±13.6; 59.94±8.3	27/26; 17/34
Joelson	2022	Sweden	Prospective cohort	DLS with LSS	slip > 3 mm	372 (decompression and fusion: 228; decompression only: 144)	63.9±8.9; 69.4±9.4	79/149; 47/97
Georgiou	2022	NSA	Retrospective cohort	DLS	single-level DLS	51619 (anterior fusion: 14971; poste- rior fusion: 36648)	≥75:1229;2514	5629/9342; 12904/23744
Chan	2021	NSA	Retrospective cohort	DLS	NA	15658	62.5	5762/9896
Takaoka	2021	Japan	Retrospective cohort	DLS with LSS	NA	145 (OLIF: 66; TLIF: 79)	66±12;71±9	28/38; 37/42
Sugiura	2021	Japan	Retrospective cohort	DLS	slippage at L3 or L4 of > 3%	202 (BPL: 51; PLIF: 106)	70.4±8.4;69.1±7.7	21/30; 38/68
Rabah	2021	USA	Retrospective cohort	DLS	NA	6260	60.27±12.49	2341/3919
Mimura	2021	Japan	Retrospective cohort	DLS	grade 1 and 2	41	69.8±7.1	19/22
Katuch	2021	Slovakia	Prospective cohort	DLS	NA	333 (TLIF: 119; PLIF:214)	55.21 ± 9.22; 56.51 ± 10.71	52/67; 95/119
Joelson	2021	Sweden	Prospective cohort	DLS with LSS	slip > 3 mm on preoperative radio- graphs	1935 (Decompression and fusion: 1338; decompression only: 597)	65±9.1;69±9.9	AN
Badhiwala	2021	Canada	Retrospective cohort	DLS	NA	1804 (laminectomy alone: 802; lami- nectomy plus fusion: 1002)	64.4±11.6; 62.7±12.1	305/497; 348/654
Zhong	2020	China	Retrospective cohort	DLS	one-level or two- level DLS	1100 (posterolateral fusion: 650; intervertebral fusion: 450)	72.80±11.7; 71.90±10.7	AN
Nyström	2020	Sweden	Retrospective cohort	DLS	slip > 3 mm	200	64.8	92/108
Lee	2020	Korea	Prospective cohort	DLS	NA	620 (decompressive laminectomy: 383; PLIF: 171; ALIF: 66)	КА	NA
Khan	2020	USA	Retrospective cohort	DLS	grade 1 and 2	850	58.35±13.78	397/453
Khan	2019	USA	Retrospective cohort	DLS	grade 1 and 2	141	62.28±10.7	65/76
Karsy	2020	USA	Prospective cohort	DLS	grade 1	608	< 60: 239; 60–70: 209; 71–80: 128;>80: 32	350/258

Chen *et al. BMC Surgery* (2023) 23:192

Table 1 (continued)	ntinued)							
Author	Year	Country	Study design	Patients	Definition of spondylolisthesis	Sample size	Age (years)	Male/female
Chan	2020	USA	Prospective cohort	DLS	grade 1	297 (MIS-TLIF: 72; Open TLIF: 225)	62.1±10.6; 59.5±11.7	32/40; 82/143
Bisson	2020	USA	Retrospective cohort	DLS	grade 1	140 (MIS decompression: 71; Open decompression: 69)	72.264 ± 9.662; 66.913 ± 12.578	74/66; 32/39
Minamide	2019	USA	Prospective cohort	DLS with LSS	single-level DLS at L3/L4 or L4/L5	218	69.7 (47–88)	96/122
Kuo	2019	USA	Retrospective cohort	DLS with LSS	NA	601 (ULBD: 164; fusion: 437)	$68.5 \pm 9.6; 69.2 \pm 9.6$	59/105; 125/312
Kelly	2019	USA	Retrospective cohort	DLS	grade 1	119 (PLF: 49; PLF + TLIF: 70)	$68 \pm 10; 65 \pm 10$	22/27; 24/46
Chan	2019	USA	Retrospective cohort	DLS	grade 1	426 (decompression alone: 84; fusion: 342)	69.9±10.5; 60.7±11.0	43/41; 131/211
Chan (1)	2019	USA	Retrospective cohort	DLS	grade 1	143 (MIS TLIF: 72; MIS decompres- sion: 71)	62.1±10.6; 72.3±9.7	32/40; 32/39
Vorhies	2018	USA	Retrospective cohort	DLS	NA	75024 (decompression: 6712; fusion: 68312)	69; 61	2776/3936; 25,645/42667
Veresciagina	2018	Switzerland	Prospective cohort	DLS	grade 1 and 2	36	66.53 (47–80)	6/30
Irmola	2018	Finland	Prospective cohort	DLS	NA	189	62 (24–87)	NA
Hayashi	2018	Japan	Retrospective cohort	DLS	grade 1 and 2	50 (CBT-PLIF: 20; MEL: 30)	69.3; 71.2	5/15; 7/23
Kato	2017	Japan	Prospective cohort	DLS	grade≤1	51	70.8±7.9	20/31
Gerling	2017	USA	Prospective cohort	DLS	ΨZ	406 (292 underwent instrumented fusion, 85 non-instrumented fusion, and 29 decompression alone)	65.3	128/278
Cheung	2016	China	Retrospective cohort	DLS	grade 1	64	69.1±8.7	29/36
Sato	2015	Japan	Retrospective cohort	DLS	NA	163 (decompression alone:74; decompression and fusion: 89)	65.8±8.9	73/90
Macki	2015	USA	Retrospective cohort	DLS	grade 1 and 2	103 (PLF:58; PLF + PLIF/TLIF:45)	59.27±11.35;55.44±10.18	26/32; 14/31
Blumenthal	2013	USA	Prospective cohort	DLS with LSS	grade 1	40	68.2±7.7	10/30
Rihn	2012	USA	Retrospective cohort	DLS	NA	601	66	189/412
Booth	1999	NSA	Retrospective cohort	DLS	NA	41	66.7 (52.2–78.7)	12/29

Author	BMI (kg/m²)	Disease duration (m)	(m) Surgery types	Follow-up (years)	Number of reoperations	Reasons for reoperation	Reoperation methods	Risk factors of reoperation
Salimi	24.3±3.9	36.2±39.6	bilateral decompres- sion	2	7	progression of lumbar degeneration or post- operative instability	NA	ЧN
hon	ΥN	NA	OD followed by spinal fusion, laminectomy, and PELD	10	549; 33	ΥZ	NA	NA
Moayeri	27.7±4.2	NA	a unilateral lami- notomy for ipsilateral decompression	8.2	22	ΥZ	decompression (12), fusion (10)	NA
Liang	23.70±3.5; 24.30±2.9	<6 mos:10;>6 mos: 94	PMTD; MIS TLIF	2	3; 1	NA	NA	NA
Joelson	27.8 ± 5.3; 26.7 ± 4.0	A	decompression and fusion; decompres- sion only	7.6	2;6	spinal stenosis or disk herniation at the index level (1.3-4), the first cranial adjacent level (L2-3), and the first caudal adjacent level (L4-5)	NA	٩N
Georgiou	ЧV	NA	anterior fusion; poste- rior fusion	2	2536; 8101	NA	NA	AN
Chan	30.7	A	posterior lumbar spine decompression with or without single level posterior instru- mented fusion	NA	438	ΥZ	NA	age, sex, BMI, smoking, addi- tion of fusion
Takaoka	NA	NA	single-level OLIF and single-level TLIF	£	3;2	NA	NA	AN
Sugiura	ΑN	NA	BPL and PLIF	m	4;4	root radiculopathy (4); ASD (3), left L5/S foraminal stenosis (1)	PLIF (3), BPL (1); PLIF (3), herniotomy (1)	AN
Rabah	30.85 ± 6.48	Å	posterior/transforami- nal lumbar interbody fusion (P/TLIF)	NA	NA	¥ Z	A	operative time, obesity class III, non-insulin- dependent dia- betes, smoking
Mimura	Ч	NA	single-level unistru- mented PLF	5	,	ASD	NA	AN
Katuch	25.56±5.12; 23.78±4.78	NA	TLIF and PLIF	S	2;15	wound infection and dural tear	NA	ЧA
Joelson	27 ± 4.5; 27 ± 4.1	NA	decompression and fusion; decompres- sion only	7.8	216; 80	spinal stenosis with or without concomitant DLS or disk herniation	ЧЧ	NA

Chen *et al. BMC Surgery* (2023) 23:192

Page 6 of 16

ladie I (continuea)	(ma							
Author	BMI (kg/m²)	Disease duration (m) Surgery types	Surgery types	Follow-up (years)	Number of reoperations	Reasons for reoperation	Reoperation methods	Risk factors of reoperation
Badhiwala	NA	AA	laminectomy alone; laminectomy plus fusion	ى ا	18, 28	related to the princi- pal procedure	NA	NA
Zhong	ΨZ	Ϋ́Α	Posterolateral fusion; intervertebral fusion	4.2	24; 9	AA	AA	BMI, sex, diabetes mellitus, age, fusion method, more bleeding
Nyström	ΝA	NA	bilateral laminotomy	6.8	29	NA	decompression (24); fusion (5)	NA
lee	NA	AA	decompressive lami- nectomy; PLIF; ALIF	10	52; 13; 1	occurrence of any lumbar spinal surgery with a disease code	NA	NA
Khan	30.96±6.21	39.64	elective open pos- terior lumbar spinal fusion	1.9	31	Ϋ́Α	NA	diabetes
Khan	30.95 ± 5.82	NA	an open posterior Iumbar fusion	1.3	7	NA	NA	NA
Karsy	Υ	AA	percutaneous screw placement, interbody placement, or decom- pression	m	38	NA	NA	A
Chan	29.5 ± 5.1; 31.3 ± 7.0	AA	minimally invasive or open TLIF	2	1;16	ASD; surgical site infection; implant removal/revision; pseudoarthrosis	NA	A
Bisson	28.735 ± 5.380; 28.190 ± 4.688	AA	MIS decompression or open decompression	m	10; 3	re-emergence of symptoms; persistent symptoms; recurrence of rt-sided pain	NA	NA
Minamide	NA	NA	micro-endoscopic decompression	2.3	17	NA	fusion (9); decompres- sion (8)	NA
Kuo	Υ Υ	NA	ULBD or fusion	3.3	17; 75	ΥZ	fusion; decompres- sion; incision and drainage, revision instrumentation, instrumentation removal	fusion as index operation, BMI, any perioperative complication, estimated blood loss
Kelly	30.6±5.9; 31.5±6.7	NA	PLF or PLF+TLIF	2	8;11	NA	NA	NA

Table 1 (continued)	led)							
Author	BMI (kg/m²)	Disease duration (m)	Surgery types	Follow-up (years)	Number of reoperations	Reasons for reoperation	Reoperation methods	Risk factors of reoperation
Chan	28.4±5.2; 31.0±6.7	<3 mos: 11;>3 mos: 401	decompression alone or fusion	-	5;15	wound revisions and/ or incision and drain- age, adjacent-seg- ment disease, screw reposition/hardware failures, bony fracture within the fusion con- struct, and evacuation of hematoma	A N	ИА
Chan (1)	29.5 ± 5.1; 28.2 ± 4.7	< 3 mos: 4;>3 mos: 133	MIS TLIF or MIS decompression	2	1;10	ASD	fusion (5); decompres- sion (6)	NA
Vorhies	NA	NA	decompression or fusion	Ŀ	686; 8469	recurrent stenosis or progressive instability	fusion; decompres- sion	NA
Veresciagina	NA	NA	laminotomy	10.78	5	NA	decompression	NA
Irmola	282±4.3	NA	instrumented lumbar spine fusion	3.9	30	acute complication; early failure; adjacent segment pathology; late failure	NA	NA
Hayashi	23.7; 23.1	39.7, 41.5	CBT-PLIF or MEL	2	3;5	a hematoma perioperatively; a vertebral fracture in the adjacent level; same-segment disease; insufficient decompression	kyphoplasty; adja- cent_x005f segment decompression; addi- tional decompression and fusion	А
Kato	ЧЧ	ΨZ	microsurgical bilateral decompression via a unilateral approach	2	7	exacerbation of disc degeneration	AN	AN
Gerling	29.2	at least 12 weeks	instrumented fusion, non-instrumented fusion, and decom- pression alone	∞	60, 20; 9	progressive spon- dylolisthesis, recurrent stenosis, complica- tions, or other	Ą	age, gender, moderate or severe stenotic levels, predomi- nant back pain, physical therapy, neurogenic clau- dication, leg pain bothersomeness scale

Author	BMI (kg/m²)	Disease duration (m) Surgery types	Surgery types	Follow-up (years)	Number of reoperations	Reasons for reoperation	Reoperation methods	Risk factors of reoperation
Cheung	₹Z	۲	decompression only	7.1	0	mechanical back pain; residual leg pain	further decompres- sion(2); Fusion(6); epidural steroid injec- tion(1)	¥ Z
Sato	23.1 ± 2.6	A	decompression alone or decompression and fusion	2	25; 13	adjacent segmen- tal disease; same segmental disease; infection; implant related; hematoma	decompression and fusion (25); decompression-only (8); others (5)	ΥN
Macki	AN	AN	NA	5.5	17; 4	degenerative disease progression	NA	NA
Blumenthal	AA	NA	laminectomy	3.6	15	mechanical low back pain	NA	NA
Rihn	29.2	at least 12 weeks	posterior decompres- sive laminectomy	4	59	Ч	NA	obesity
Booth	₹ Z	A	decompression, autogenous iliac crest bone grafting, inter- transverse process fusion, and segmental (pedicle screw) instru- mentation	6.5	Ŷ	recurrent stenosis at adjacent levels with transition syndrome	NA	A

oblique lateral interbody fusion, *TLF* transforaminal lumbar interbody fusion, *BPL* bilateral partial laminectomy, *PLL* posterior rumbar interbody fusion, *TLF* transforaminal lumbar interbody fusion, *BPL* bilateral partial laminectomy, *PLL* posterior rumbar interbody fusion with cortical bone trajectory, *MEL* microendoscopic laminotomy, *OD* open discectomy, *PELD* percutaneous endoscopic lumbar discectomy, *PMTD* paraspinal mini-tubular lumbar discectomy, *PLL* paraspinal mini-tubular bilateral paraspinal mini-tubular bilateral paraspinal mini-tubular bilateral decompression, *ASD* adjacent segment disease, *BMI* body mass index, *NA* not available

uthor (Year)	Rate (95% CI)	Weigh
Galimi (2022)	0.04 (-0.01, 0.09)	1.72
loh_a (2022)	0.14 (0.13, 0.15)	1.95
loh_b (2022)	0.13 (0.09, 0.17)	1.82
Noayeri (2022)	0.16 (0.10, 0.22)	1.68
iang_a (2022)	0.06 (-0.01, 0.12)	1.66
iang_b (2022)	0.02 (-0.02, 0.06)	1.84
oelson_a (2022)	0.01 (-0.00, 0.02)	1.95
oelson_b (2022)	0.04 (0.01, 0.07)	1.87
eorgiou_a (2022)	0.17 (0.16, 0.18)	1.96
eorgiou_b (2022)	0.22 (0.22, 0.23)	1.96
han (2021)	0.03 (0.03, 0.03)	1.96
akaoka_a (2021)	0.05 (-0.00, 0.10)	1.75
akaoka_b (2021)	0.03 (-0.01, 0.06)	1.86
ugiura_a (2021)	0.08 (0.00, 0.15)	1.56
ugiura_b (2021)	0.04 (0.00, 0.07)	1.85 1.78
limura (2021) +	0.02 (-0.02, 0.07) 0.02 (-0.01, 0.04)	1.91
atuch_a (2021)	0.02 (-0.01, 0.04)	1.8
belson_a (2021)	0.16 (0.14, 0.18)	1.92
belson_b (2021)	0.13 (0.11, 0.16)	1.89
adhiwala_a (2021)	0.02 (0.01, 0.03)	1.95
adhiwala_b (2021)	0.03 (0.02, 0.04)	1.95
yström (2020)	0.14 (0.10, 0.19)	1.76
ee_a (2020)	0.14 (0.10, 0.17)	1.86
ee_b (2020)	0.08 (0.04, 0.12)	1.83
ee_c (2020)	0.02 (-0.01, 0.04)	1.88
han (2020) 🔷 🖡	0.04 (0.02, 0.05)	1.94
han (2019) 🔶	0.05 (0.01, 0.09)	1.85
arsy (2020)	0.06 (0.04, 0.08)	1.93
han_a (2020)	0.01 (-0.01, 0.04)	1.89
han_b (2020)	0.07 (0.04, 0.10)	1.86
isson_a (2020)	0.14 (0.06, 0.22)	1.50
isson_b (2020)	0.04 (-0.00, 0.09)	1.77
linamide (2019)	0.08 (0.04, 0.11)	1.85
uo_a (2019)	0.10 (0.06, 0.15)	1.78
uo_b (2019)	0.17 (0.14, 0.21)	1.85
elly_a (2019)	0.16 (0.06, 0.27)	1.31
elly_b (2019)	0.16 (0.07, 0.24)	1.47 1.75
han_a (2019)	0.06 (0.01, 0.11) 0.04 (0.02, 0.07)	1.92
han (1) (2019)	0.14 (0.06, 0.22)	1.50
orhies_a (2018)	0.10 (0.09, 0.11)	1.95
orhies_b (2018)	0.12 (0.12, 0.13)	1.96
eresciagina (2018)	 0.14 (0.03, 0.25) 	1.23
mola (2018)	0.16 (0.11, 0.21)	1.74
ayashi_a (2018)	0.15 (-0.01, 0.31)	0.92
ayashi_b (2018)	0.17 (0.03, 0.30)	1.07
ato (2017)	0.04 (-0.01, 0.09)	1.73
erling_a (2017)	0.21 (0.16, 0.25)	1.78
erling_b (2017)	0.24 (0.15, 0.33)	1.42
erling_c (2017)	0.31 (0.14, 0.48)	0.85
heung (2016)	0.14 (0.06, 0.23)	1.47
ato_a (2015)	0.34 (0.23, 0.45)	1.27
ato_b (2015)	0.15 (0.07, 0.22)	1.57
acki_a (2015)	0.29 (0.18, 0.41)	1.20
lacki_b (2015)	0.09 (0.01, 0.17)	1.48
lumenthal (2013)	0.38 (0.22, 0.53)	0.96
ihn (2012)	0.10 (0.07, 0.12)	1.91
ooth (1999)	0.12 (0.02, 0.22)	1.34
overall, DL (l ² = 99.3%, p = 0.000)	0.10 (0.08, 0.12)	100.00
5 0	.5	

Fig. 2 Forest plot regarding to reoperation rate

(95%CI: 8%-13%). In patients subgroup, DLS patients showed 11% of reoperation rate (95%CI: 8%-13%) and DLS patients with LSS showed 10% of reoperation rate (95%CI: 6%-13%). In surgery types subgroup, the reoperation rate was 11% (95%CI: 9%-13%) in patients undergoing decompression, 10% (95%CI: 7%-12%) in patients undergoing fusion, 9% (95%CI: 5%-13%) in patients undergoing decompression and fusion, and 7% (95%CI: 3%-11%) in patients undergoing other surgeries. In follow-up time subgroup, the reoperation rate was 9% (95%CI: 6%-12%), 12% (95%CI: 9%-14%), and 10% (95%CI: 6%-15%) at follow-up time < 5 years, between 5 to 10 years, and \geq 10 years, respectively. In study quality subgroup, there was 11% (95%CI: 9%-13%) of reoperation rate in studies with fair quality and 7% (95%CI: 5%-10%) of reoperation rate in studies with high quality. The overall and subgroup analysis results were shown in Table 2.

Meta-analysis of risk factors for reoperation

The meta-analysis showed that obesity (OR=1.91, 95%CI: 1.04–3.51, I^2 =53.1%), diabetes (OR=2.01, 95%CI: 1.43–2.82, I^2 =0%), and smoking (OR=1.51, 95%CI: 1.23–1.84, I^2 =0%) were associated with an increased risk of reoperation. Age (OR=0.99, 95%CI: 0.95–1.03, I^2 =78.4%), sex (OR=1.31, 95%CI: 0.83–2.05,

Table 2 Me	ta analysis	of reoperati	on rate
------------	-------------	--------------	---------

Outcomes	Number of studies	Rate (95%CI)	l ²
Reoperation rate	37	0.10 (0.08–0.12)	99.3
Sensitivity analysis		0.10 (0.08–0.12)	
Publication bias		Z=2.91	P = 0.004
Study design			
Prospective cohort	12	0.10 (0.07–0.13)	94.5
Retrospective cohort	25	0.10 (0.08–0.13)	99.5
Patients			
DLS	27	0.11 (0.08–0.13)	99.4
DLS with LSS	10	0.10 (0.06–0.13)	96.5
Surgery types			
Decompression	21	0.11 (0.09–0.13)	91.9
Fusion	8	0.10 (0.07–0.12)	98.9
Decompression and fusion	6	0.09 (0.05–0.13)	99.0
Others	2	0.07 (0.03–0.11)	23.4
Follow-up (years)			
<5	20	0.09 (0.06–0.12)	98.5
5–10	13	0.12 (0.09–0.14)	98.1
≥10	3	0.10 (0.06–0.15)	92.8
Quality			
Fair	26	0.11 (0.09–0.13)	98.6
High	11	0.07 (0.05–0.10)	96.7

Abbreviation: CI confidence interval, I² I-squared, DLS degenerative lumbar spondylolisthesis, LSS lumbar spinal stenosis

 I^2 =60.4%), and more bleeding (OR=0.86, 95%CI: 0.07–10.22, I^2 =87.5%) were not associated with the reoperation. The overall results were demonstrated in Table 3. Forest plots regarding to obesity, diabetes, and smoking were demonstrated in Fig. 3A, B, and C, respectively.

Systematic review of risk factors for reoperation

This systematic review examined two literatures about the obesity. Chan et al. carried out a retrospective cohort study of obesity and reoperation after lumbar surgery [23]. As expected, significant higher risk of reoperation was found in patients who were obese [23]. Similar evidence was supported by Rabah et al. that an increase of one unit in BMI was associated with 4% increased risk of reoperation [11]. Moreover, study of Chan et al. showed addition of fusion was associated with higher risk of reoperation [23]. Rabah et al. found operative time > 5 h to be associated with an increased risk of reoperation [11]. A study consisted of 5-year follow-up indicated that having an index of fusion operation and perioperative complications was associated with the increased odds of reoperation [13]. Compared to intervertebral fusion, patients undergoing posterolateral fusion had 4.02-times risk of reoperation [8]. In addition, Gerling et al. have reported that patients with 2/3 moderate or severe stenotic levels, predominant back pain, no physical therapy, and greater leg pain score at baseline indicated higher reoperation rate [26].

Assessment of publication bias and sensitivity analysis

Sensitivity analysis was performed by sequentially removing the study to assess the robustness of overall results. All results of sensitivity analysis were consistent with those of the main analysis (Tables 2 and 3). By

 Table 3
 Risk factors for the reoperation of DLS patients after surgeries

Risk factors	Number of studies	OR (95%CI)	Ρ	l ²
Age	3	0.99 (0.95–1.03)	0.535	78.4
Sensitivity analysis		0.99 (0.95–1.03)		
Sex	3	1.31 (0.83–2.05)	0.243	60.4
Sensitivity analysis		1.31 (0.83–2.05)		
Obesity	3	1.91 (1.04–3.51)	0.037	53.1
Sensitivity analysis		1.91 (1.04–3.51)		
Diabetes	3	2.01 (1.43–2.82)	< 0.001	0.0
Sensitivity analysis		2.01 (1.43–2.82)		
Smoking	2	1.51 (1.23–1.84)	< 0.001	0.0
Sensitivity analysis		1.51 (1.23–1.84)		
More bleeding	2	0.86 (0.07–10.22)	0.903	87.5
Sensitivity analysis		0.86 (0.07–10.22)		

Abbreviation: DLS degenerative lumbar spondylolisthesis, OR odds ratio, Cl confidence interval, l^2 I-squared

Author (Year)		OR (95% CI)	% Weight
Zhong (2020)		- 4.42 (1.50, 14.06)	20.22
Kuo (2019) —	-	1.19 (0.64, 2.23)	38.23
Rihn (2012)		1.97 (1.13, 3.44)	41.55
Overall, DL (l ² = 53.1%, p = 0.119)		1.91 (1.04, 3.51)	100.00
.0625	1	16	
NOTE: Weights are from random-effects model			

В				
				%
	Author (Year)		OR (95% CI)	Weight
	Rabah (2021)		1.85 (1.25, 2.69)	77.98
	Zhong (2020) —		3.08 (0.78, 7.85)	8.59
	Khan (2020)		2.49 (0.99, 6.27)	13.43
	Overall, IV (I ² = 0.0%, p = 0.634)	$\langle \rangle$	2.01 (1.43, 2.82)	100.00
	.125	1	3	

C			
			%
Author (Year)		OR (95% CI)	Weight
Chan (2021)		1.41 (1.10, 1.80)	66.19
Rabah (2021)		— 1.72 (1.21, 2.41)	33.81
Overall, IV (l ² = 0.0%, p = 0.358)		1.51 (1.23, 1.84)	100.00
	1 2		

Fig. 3 Forest plots regarding to obesity (A), diabetes (B), and smoking (C)

funnel plot, we detected an evidence of publication biases (Z=2.91, P=0.004) (Table 2, Supplementary Fig. 1A). Therefore, a trim-and-fill method was utilized to fill the missing data to eliminate the impact of publication bias. Funnel plot with missing data filled was demonstrated in Supplementary Fig. 1B. Before filled, reoperation rate was 10% (95%CI: 8%-12%). After filled, reoperation rate was 11% (95%CI: 9%-13%).

Discussion

The reoperation rate of DLS patients undergoing lumbar surgeries remains high in spite of improved surgical skills and techniques; therefore, exploring risk factors of reoperation is important [7, 8]. Considering the controversial results in the risk factors [8, 11–13], we performed a systematic review and meta-analysis based on currently available studies to analyze the reoperation rate and risk factors. In this study, we found a 10% of reoperation rate in DLS patients after lumbar surgeries. Obesity, diabetes, and smoking were identified as risk factors for the reoperation.

Several previous studies have reported the reoperation rate after lumbar surgeries in DLS patients [7, 49-52]. The reoperation rate was reported as 12.4% from 1990 to 1993 and 14.0% from 1997 to 2000 [51]. Ghogawala et al. proved that the reoperation rate was 15% at 1 year after the surgery in DLS patients only undergoing decompression [52]. In the present studies, the reoperation rate was found nearly the same as that reported in previous studies [7]. The reoperation rate in DLS patients was 15.7% at the mean follow-up of 8.2 years [7]. For patients undergoing fusion procedures, the cumulative reoperation rate was 14% [49]. Another report demonstrated that the reoperation rate ranged from 5.8% to 16.3% according to the type of surgeries [50]. Similar to the studies mentioned above, in our study, the reoperation rate of DLS was 10%, ranging from 8 to 12%. Our results may be useful for clinicians to evaluate the reoperation rate.

Identifying risk factors of reoperation for patients after lumbar surgeries is of clinical interest. In this study, obesity, diabetes, and smoking were found to be associated with higher risk of reoperation. Rabah et al. and Chan et al. have confirmed that smoking status was associated with greater risk of reoperation [11, 23]. Also, there were several studies reporting the positive association between obesity and reoperation of patients undergoing lumbar surgeries [8, 23]. This can be explained by that obese patients were more likely to be frail [53, 54], and frail patients had 56% increased odds of reoperation after lumbar surgery [23].

Animal studies have long recognized the close association between diabetes and lumbar spine disorders [55– 57]. Diabetic models have revealed some harmful changes, such as increase of toxic end products of glycation, expression of matrix metalloproteinases 2 related to extracellular matrix degradation, and hyperglycemia-induced intervertebral disc inflammation, promoting intervertebral disc degeneration process [58-60]. Studies have revealed that diabetes was closely associated with degenerative lumbar spine disorders [61, 62]. Park et al. have found the influence of diabetes on the prevalence of lumbar spine surgeries, indicating that diabetes may be a factor aggravating lumbar spine disorders [62]. In Park et al. study, patients with diabetes underwent more lumbar surgeries than those without diabetes [62]. Their finding suggested that diabetes was significantly associated with the increased number of lumbar spine surgeries, and this finding is of critical importance because it revealed that diabetes may be an incentive for the increase of the severity of lumbar spine disorders, which ultimately led to the necessity of surgeries [62]. In this meta-analysis, diabetes was identified as a risk factor for the reoperation of DLS patients undergoing lumbar surgeries. This was consistent with the findings from Zhong et al. [8] Our findings suggested that when treating DLS patients with diabetes, physicians should pay more attention to glycemic control for the purpose of decreasing the risk of reoperation.

This meta-analysis explores the reoperation rate and risk factors for the reoperation. Results show that there is 10% of reoperation after lumbar surgeries, and obesity, diabetes, and smoking are found to increase the risk of reoperation. Our findings suggest that DLS patients should control glycemic level and weight, and reduce smoking to decrease the risk of reoperation. There are some limitations in this study. First, all fusion techniques (PLF and LIF) were put together. Due to the limitations of the included studies, it is unable to further analyze the reoperation rate in DLS patients undergoing the single fusion technique. Second, the number of studies reporting the risk factors of reoperation is relatively small, and some outcomes can only be qualitatively described, which may affect the stability of the results. Third, the risk of reoperation may be different according to the severity of lumbar spondylolisthesis and the first surgical methods; however, data provided in the currently available studies are insufficient to further analyze. Future meta-analysis including more relevant studies are needed to verify our findings and to explore the effect of lumbar spondylolisthesis severity and the first surgical methods on the risk of reoperation.

Conclusion

Our meta-analysis found 10% of reoperation rate in DLS patients undergoing lumbar surgeries, and identified obesity, diabetes, and smoking as risk factors for the reoperation. Our findings suggested that patients should improve glycemic level and weight, and quit smoking to reduce the reoperation after lumbar surgery.

Abbreviations

DLS	Degenerative lumbar spondylolisthesis
PRISMA	Preferred Reporting Items for Systematic Reviews and
	Meta-Analyses
BMI	Body mass index
NOS	Newcastle-Ottawa Scale
OR	Odds ratio
CI	Confidence interval

Supplementary Information

The online version contains supplementary material available at https://doi.org/10.1186/s12893-023-02082-8.

Additional file 1: Supplementary figure 1. Funnel plot for publication bias before and after trim-and-fillanalysis.

Acknowledgements

Not applicable.

Authors' contributions

YC and XF designed the study. YC wrote the manuscript. YZ, JC, YL and YW collected, analyzed and interpreted the data. XF critically reviewed, edited and approved the manuscript. All authors read and approved the final manuscript.

Funding

This study was funded by Sichuan Province Key Research and Development Project (2022YFS0418).

Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Declarations

Ethics approval and consent to participate Not applicable.

Consent for publication

Not applicable.

Competing interests

The authors declare no competing interests.

Received: 6 February 2023 Accepted: 16 June 2023 Published online: 05 July 2023

References

- Bydon M, Alvi MA, Goyal A. Degenerative lumbar spondylolisthesis: definition, natural history, conservative management, and surgical treatment. Neurosurg Clin N Am. 2019;30:299–304.
- Wang YXJ, Káplár Z, Deng M, Leung JCS. Lumbar degenerative spondylolisthesis epidemiology: a systematic review with a focus on genderspecific and age-specific prevalence. J Orthop Translat. 2017;11:39–52.
- Ravindra VM, Senglaub SS, Rattani A, Dewan MC, Härtl R, Bisson E, et al. Degenerative lumbar spine disease: estimating global incidence and worldwide volume. Global spine journal. 2018;8:784–94.
- Karsy M, Bisson EF. Surgical versus nonsurgical treatment of lumbar spondylolisthesis. Neurosurg Clin N Am. 2019;30:333–40.
- Kim CH, Chung CK, Choi Y, Kim MJ, Kim MJ, Shin S, et al. Increased proportion of fusion surgery for degenerative lumbar spondylolisthesis and changes in reoperation rate: a nationwide cohort study with a minimum 5-year follow-up. Spine. 2019;44:346–54.
- Gaderer C, Schaumann A, Schulz M, Thomale UW. Neuroendoscopic lavage for the treatment of CSF infection with hydrocephalus in children. Child's Nerv Syst. 2018;34:1893–903.

- Moayeri N, Rampersaud YR. Revision surgery following minimally invasive decompression for lumbar spinal stenosis with and without stable degenerative spondylolisthesis: a 5- to 15-year reoperation survival analysis. J Neurosurg Spine. 2021;36:1–7.
- Zhong W, Liang X, Luo X, Huang T, Quan Z. Complications rate of and risk factors for the unplanned reoperation of degenerative lumbar spondylolisthesis in elderly patients: a retrospective single-Centre cohort study of 33 patients. BMC Geriatr. 2020;20:301.
- Park MS, Ju YS, Moon SH, Kim TH, Oh JK, Sung PS, et al. Reoperation rates after posterior lumbar spinal fusion surgery according to preoperative diagnoses: a national population-based cohort study. Clin Neurol Neurosurg. 2019;184:105408.
- Noh SH, Cho PG, Kim KN, Lee B, Lee JK, Kim SH. Risk factors for reoperation after lumbar spine surgery in a 10-year Korean national health insurance service health examinee cohort. Sci Rep. 2022;12:4606.
- Rabah NM, Khan HA, Shost M, Beckett J, Mroz TE, Steinmetz MP. Predictors of operative duration and complications in single-level posterior interbody fusions for degenerative spondylolisthesis. World neurosurgery. 2021;151:e317–23.
- Khan JM, Michalski J, Basques BA, Louie PK, Chen O, Hayani Z, et al. Do Clinical outcomes and sagittal parameters differ between diabetics and nondiabetics for degenerative spondylolisthesis undergoing lumbar fusion? Global spine journal. 2020;10:286–93.
- Kuo CC, Merchant M, Kardile MP, Yacob A, Majid K, Bains RS. In Degenerative Spondylolisthesis, Unilateral Laminotomy for Bilateral Decompression Leads to Less Reoperations at 5 Years When Compared to Posterior Decompression With Instrumented Fusion: a propensity-matched retrospective analysis. Spine. 2019;44:1530–7.
- Moher D, Liberati A, Tetzlaff J, Altman DG. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. Ann Intern Med. 2009;151(264–9):w64.
- Xue X, Lu CL, Jin XY, Liu XH, Yang M, Wang XQ, et al. Relationship between serum uric acid, all-cause mortality and cardiovascular mortality in peritoneal dialysis patients: systematic review and meta-analysis of cohort studies. BMJ Open. 2021;11:e052274.
- Badhiwala JH, Leung SN, Jiang F, Wilson JRF, Akbar MA, Nassiri F, et al. In-hospital course and complications of laminectomy alone versus laminectomy plus instrumented posterolateral fusion for lumbar degenerative spondylolisthesis: a retrospective analysis of 1804 patients from the NSQIP Database. Spine. 2021;46:617–23.
- Bisson EF, Mummaneni PV, Virk MS, Knightly J, Alvi MA, Goyal A, et al. Open versus minimally invasive decompression for low-grade spondylolisthesis: analysis from the Quality Outcomes Database. J neurosurg Spine. 33(3):349-59.
- Blumenthal C, Curran J, Benzel EC, Potter R, Magge SN, Harrington JF Jr, et al. Radiographic predictors of delayed instability following decompression without fusion for degenerative grade I lumbar spondylolisthesis. J Neurosurg Spine. 2013;18:340–6.
- Booth KC, Bridwell KH, Eisenberg BA, Baldus CR, Lenke LG. Minimum 5-year results of degenerative spondylolisthesis treated with decompression and instrumented posterior fusion. Spine. 1999;24:1721–7.
- Chan AK, Bisson EF, Bydon M, Glassman SD, Foley KT, Potts EA, et al. A comparison of minimally invasive transforaminal lumbar interbody fusion and decompression alone for degenerative lumbar spondylolisthesis. Neurosurg Focus. 2019;46:E13.
- Chan AK, Bisson EF, Bydon M, Glassman SD, Foley KT, Potts EA, et al. Laminectomy alone versus fusion for grade 1 lumbar spondylolisthesis in 426 patients from the prospective Quality Outcomes Database. J Neurosurg Spine. 2018;30:234–41.
- Chan AK, Bisson EF, Bydon M, Foley KT, Glassman SD, Shaffrey CI, et al. A Comparison of Minimally Invasive and Open Transforaminal Lumbar Interbody Fusion for Grade 1 Degenerative Lumbar Spondylolisthesis: An Analysis of the Prospective Quality Outcomes Database. Neurosurgery. 2020;87:555–62.
- Chan V, Witiw CD, Wilson JRF, Wilson JR, Coyte P, Fehlings MG. Frailty is an important predictor of 30-day morbidity in patients treated for lumbar spondylolisthesis using a posterior surgical approach. Spine J. 2022;22:286–95.
- Cheung JP, Cheung PW, Cheung KM, Luk KD. Decompression without Fusion for Low-Grade Degenerative Spondylolisthesis. Asian Spine J. 2016;10:75–84.

- Georgiou S, Saggi S, Wu HH, Metz L. Comparison of 90-day complications and two-year reoperation rates between anterior and posterior interbody fusion for single-level degenerative spondylolisthesis. N Am Spine Soc J. 2022;10:100127.
- Gerling MC, Leven D, Passias PG, Lafage V, Bianco K, Lee A, et al. Risk Factors for Reoperation in Patients Treated Surgically for Degenerative Spondylolisthesis: a subanalysis of the 8-year data from the SPORT Trial. Spine. 2017;42:1559–69.
- Hayashi K, Toyoda H, Terai H, Hoshino M, Suzuki A, Takahashi S, et al. Comparison of minimally invasive decompression and combined minimally invasive decompression and fusion in patients with degenerative spondylolisthesis with instability. J Clin Neurosci. 2018;57:79–85.
- Irmola TM, Häkkinen A, Järvenpää S, Marttinen I, Vihtonen K, Neva M. Reoperation Rates Following Instrumented Lumbar Spine Fusion. Spine. 2018;43:295–301.
- Joelson A, Nerelius F, Holy M, Sigmundsson FG. Reoperations after decompression with or without fusion for L4–5 spinal stenosis with or without degenerative spondylolisthesis: a study of 6,532 patients in Swespine, the national Swedish spine register. Acta Orthop. 2021;92:264–8.
- Joelson A, Nerelius F, Holy M, Sigmundsson FG. Reoperations After Decompression With or Without Fusion for L3–4 Spinal Stenosis With Degenerative Spondylolisthesis: A Study of 372 Patients in Swespine, the National Swedish Spine Register. Clini Spine Surg. 2022;35:E389–93.
- Karsy M, Chan AK, Mummaneni PV, Virk MS, Bydon M, Glassman SD, et al. Outcomes and complications with age in spondylolisthesis: an evaluation of the elderly from the quality outcomes database. Spine. 2020;45:1000–8.
- Kato M, Namikawa T, Matsumura A, Konishi S, Nakamura H. Radiographic risk factors of reoperation following minimally invasive decompression for lumbar canal stenosis associated with degenerative scoliosis and spondylolisthesis. Global Spine J. 2017;7:498–505.
- Katuch V, Grega R, Knorovsky K, Banoci J, Katuchova J, Sasala M, et al. Comparison between posterior lumbar interbody fusion and transforaminal lumbar interbody fusion in the management of lumbar spondylolisthesis. Bratisl Lek Listy. 2021;122:653–6.
- Kelly JP, Alcala-Marquez C, Dawson JM, Mehbod AA, Pinto MR. Treatment of degenerative spondylolisthesis by instrumented posterolateral versus instrumented posterolateral with transforaminal lumbar interbody singlelevel fusion. J Spine Surg (Hong Kong). 2019;5:351–7.
- 35. Khan JM, Harada GK, Basques BA, Nolte MT, Louie PK, Iloanya M, et al. Patients with predominantly back pain at the time of lumbar fusion for low-grade spondylolisthesis experience similar clinical improvement to patients with predominantly leg pain: mid-term results. Spine J. 2020;20:276–82.
- Lee CH, Kim CH, Chung CK, Choi Y, Kim MJ, Yim D, et al. Long-term effect of diabetes on reoperation after lumbar spinal surgery: a nationwide population-based sample cohort study. World Neurosurg. 2020;139:e439–48.
- Liang Z, Xu X, Rao J, Chen Y, Wang R, Chen C. Clinical evaluation of Paraspinal mini-tubular lumbar decompression and minimally invasive Transforaminal lumbar interbody fusion for lumbar spondylolisthesis grade i with lumbar spinal stenosis: a cohort study. Front Surg. 2022;9:906289.
- Macki M, Bydon M, Weingart R, Sciubba D, Wolinsky JP, Gokaslan ZL, et al. Posterolateral fusion with interbody for lumbar spondylolisthesis is associated with less repeat surgery than posterolateral fusion alone. Clin Neurol Neurosurg. 2015;138:117–23.
- Mimura T, Tsutsumimoto T, Yui M, Misawa H. Does fusion status following posterolateral lumbar fusion in the treatment for stable lumbar degenerative spondylolisthesis affect the long-term surgical outcomes? A propensity score-weighted analysis of consecutive patients. J Orthop Sci. 2022;27:990–4.
- Minamide A, Simpson AK, Okada M, Enyo Y, Nakagawa Y, Iwasaki H, et al. Microendoscopic Decompression for Lumbar Spinal Stenosis With Degenerative Spondylolisthesis: The Influence of Spondylolisthesis Stage (Disc Height and Static and Dynamic Translation) on Clinical Outcomes. Clin Spine Surg. 2019;32:E20–6.

- Nyström B, Jin S, Schillberg B, Moström U, Lundin P, Taube A. Are degenerative spondylolisthesis and further slippage postoperatively really issues in spinal stenosis surgery? Scand J Pain. 2020;20:307–17.
- 42. Rihn JA, Radcliff K, Hilibrand AS, Anderson DT, Zhao W, Lurie J, et al. Does obesity affect outcomes of treatment for lumbar stenosis and degenerative spondylolisthesis? Analysis of the Spine Patient Outcomes Research Trial (SPORT). Spine. 2012;37:1933–46.
- 43. Salimi H, Toyoda H, Terai H, Yamada K, Hoshino M, Suzuki A, et al. Mid-term changes in spinopelvic sagittal alignment in lumbar spinal stenosis with coexisting degenerative spondylolisthesis or scoliosis after minimally invasive lumbar decompression surgery: minimum five-year follow-up. Spine J. 2022;22:819–26.
- Sato S, Yagi M, Machida M, Yasuda A, Konomi T, Miyake A, et al. Reoperation rate and risk factors of elective spinal surgery for degenerative spondylolisthesis: minimum 5-year follow-up. Spine J. 2015;15:1536–44.
- Sugiura T, Okuda S, Takenaka S, Nagamoto Y, Matsumoto T, Takahashi Y, et al. Comparing Investigation Between Bilateral Partial Laminectomy and Posterior Lumbar Interbody Fusion for Mild Degenerative Spondylolisthesis. Clin Spine Surg. 2021;34:E403–9.
- Takaoka H, Inage K, Eguchi Y, Shiga Y, Furuya T, Maki S, et al. Comparison between intervertebral oblique lumbar interbody fusion and transforaminal lumbar interbody fusion: a multicenter study. Sci Rep. 2021;11:16673.
- Veresciagina K, Mehrkens A, Schären S, Jeanneret B. Minimum ten-year follow-up of spinal stenosis with degenerative spondylolisthesis treated with decompression and dynamic stabilization. J Spine Surg (Hong Kong). 2018;4:93–101.
- Vorhies JS, Hernandez-Boussard T, Alamin T. Treatment of Degenerative Lumbar Spondylolisthesis With Fusion or Decompression Alone Results in Similar Rates of Reoperation at 5 Years. Clin Spine Surg. 2018;31:E74–9.
- Ghogawala Z, Dziura J, Butler WE, Dai F, Terrin N, Magge SN, et al. Laminectomy plus Fusion versus Laminectomy Alone for Lumbar Spondylolisthesis. N Engl J Med. 2016;374:1424–34.
- Schöller K, Alimi M, Cong GT, Christos P, Härtl R. Lumbar Spinal Stenosis Associated With Degenerative Lumbar Spondylolisthesis: A Systematic Review and Meta-analysis of Secondary Fusion Rates Following Open vs Minimally Invasive Decompression. Neurosurgery. 2017;80:355–67.
- Mardjetko SM, Connolly PJ, Shott S. Degenerative lumbar spondylolisthesis. A meta-analysis of literature 1970–1993. Spine. 1994;19:2256s–65s.
- Ghogawala Z, Benzel EC, Amin-Hanjani S, Barker FG 2nd, Harrington JF, Magge SN, et al. Prospective outcomes evaluation after decompression with or without instrumented fusion for lumbar stenosis and degenerative Grade I spondylolisthesis. J Neurosurg Spine. 2004;1:267–72.
- Liao Q, Zheng Z, Xiu S, Chan P. Waist circumference is a better predictor of risk for frailty than BMI in the community-dwelling elderly in Beijing. Aging Clin Exp Res. 2018;30:1319–25.
- Rietman ML, van der AD, van Oostrom SH, Picavet HSJ, Dollé MET, van Steeg H, et al. The Association between BMI and Different Frailty Domains: A U-Shaped Curve? J Nutr Health Aging. 2018;22:8–15.
- Fields AJ, Berg-Johansen B, Metz LN, Miller S, La B, Liebenberg EC, et al. Alterations in intervertebral disc composition, matrix homeostasis and biomechanical behavior in the UCD-T2DM rat model of type 2 diabetes. J Orthop Res. 2015;33:738–46.
- Illien-Junger S, Grosjean F, Laudier DM, Vlassara H, Striker GE, latridis JC. Combined anti-inflammatory and anti-AGE drug treatments have a protective effect on intervertebral discs in mice with diabetes. PLoS One. 2013;8:e64302.
- Illien-Jünger S, Lu Y, Qureshi SA, Hecht AC, Cai W, Vlassara H, et al. Chronic ingestion of advanced glycation end products induces degenerative spinal changes and hypertrophy in aging pre-diabetic mice. PLoS One. 2015;10:e0116625.
- Cheng X, Ni B, Zhang Z, Liu Q, Wang L, Ding Y, et al. Polyol pathway mediates enhanced degradation of extracellular matrix via p38 MAPK activation in intervertebral disc of diabetic rats. Connect Tissue Res. 2013;54:118–22.
- Chen S, Liao M, Li J, Peng H, Xiong M. The correlation between microvessel pathological changes of the endplate and degeneration of the intervertebral disc in diabetic rats. Exp Ther Med. 2013;5:711–7.

- Won HY, Park JB, Park EY, Riew KD. Effect of hyperglycemia on apoptosis of notochordal cells and intervertebral disc degeneration in diabetic rats. J Neurosurg Spine. 2009;11:741–8.
- Agius R, Galea R, Fava S. Bone mineral density and intervertebral disc height in type 2 diabetes. J Diabetes Complications. 2016;30:644–50.
- Park CH, Min KB, Min JY, Kim DH, Seo KM, Kim DK. Strong association of type 2 diabetes with degenerative lumbar spine disorders. Sci Rep. 2021;11:16472.

Publisher's Note

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Ready to submit your research? Choose BMC and benefit from:

- fast, convenient online submission
- thorough peer review by experienced researchers in your field
- rapid publication on acceptance
- support for research data, including large and complex data types
- gold Open Access which fosters wider collaboration and increased citations
- maximum visibility for your research: over 100M website views per year

At BMC, research is always in progress.

Learn more biomedcentral.com/submissions

