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

Anders JOELSON, Fredrik NERELIUS, Marek HOLY, and Freyr Gauti SIGMUNDSSON

Department of Orthopedics, Örebro University School of Medical Sciences and Örebro University Hospital, Örebro, Sweden
Correspondence: anders@joelson.se
Submitted 2020-09-19. Accepted 2020-11-16.

There is an ongoing controversy on how to surgically address lumbar spinal stenosis with and without concomitant degenerative spondylolisthesis (DS). The debate has focused on the durability of the index operation versus accelerated symptomatic degeneration at the adjacent segments. In this debate, durability has been defined as maintenance of clinical benefit without the need for additional intervention (Ghogawala et al. 2017). While decompression without fusion may increase the risk for early index-level reoperations for instability (Ghogawala et al. 2016, Urakawa et al. 2020), fusion surgery might increase the risk of adjacent segment disease (ASD) (Sears et al. 2011, Okuda et al. 2018).

The national Swedish spine register (Swespine) covers 90% of the spine units in Sweden and the follow-up rate is 75–80% (Strömqvist et al. 2013). The Swespine offers possibilities to examine outcome in a large dataset of patients operated on for isolated L4–5 disease, which is the most common clinical scenario within spine surgery. Large register studies may contribute to increased evidence in this area.

Using Swespine data, we investigated reoperation rates at the index and adjacent levels after L4–5 fusion surgery in a large cohort of unselected patients registered in Swespine, the national Swedish spine register.

Patients and methods
Study design
National Register study with prospectively collected data.

Patients
The Swespine is a national quality register administered by the Swedish Association of Spine Surgeons. Demographic information is registered and follow-up forms are mailed to patients with a prepaid return envelope. Participation is volun-
We identified 6,584 patients who underwent surgery for lumbar L4–5 spinal stenosis with or without DS between 2007 and 2012. 52 patients were subject to acute reoperations (e.g., due to postoperative hematomas) and were excluded. The remaining 6,532 patients were followed up between 2007 and 2017 to identify reoperations. The diagnosis spinal stenosis was reported by the operating surgeon. DS was defined as slip > 3 mm on preoperative radiographs. Only patients who underwent surgery at the L4–5 level were included in the study.

Reoperation rates

In Swespine, index operations and reoperations are registered separately. In order to identify reoperations that were incorrectly classified as index operations, we scanned the register for patients who had multiple operations. Reoperations because of spinal stenosis with or without concomitant DS or disk herniation at the index level (L4–5), the 1st cranial adjacent level (L3–4), the 2nd cranial adjacent level (L2–3), and the 1st caudal adjacent level (L5–S1) were counted. The diagnoses spinal stenosis and disk herniation were reported by the operating surgeon. Only the 1st reoperation was counted. Reoperations at the index level (L4–5) because of implant failure were counted separately.

Patient-reported outcome measures

Patient-reported outcome measures (PROMs) were recorded preoperatively, and at 1 and 2 years after surgery. We evaluated pain, disability, and health-related quality of life. Numeric rating scales (NRS) were used to assess leg and back pain (range 0–10, 0 being the best). The Oswestry disability index version 2.0 (ODI) (Fairbank and Pynsent 2000) was used to assess disability (range 0–100, 0 being the best and 100 the worst). The EQ5D index (UK tariff) (EuroQol Group 1990, Dolan 1997) was used to assess health-related quality of life (range –0.59 to 1, –0.59 being the worst and 1 the best).

Statistics

The results are presented as mean (SD). Student’s t-test for unpaired data was used to compare normally distributed data. The Mann–Whitney U-test was used to compare unpaired non-normally distributed data. The chi-square test was used to compare frequencies. A p-value of < 0.05 was considered significant, and 2-tailed tests were used. The 95% confidence interval (CI) for relative risks is calculated as described by Altman (1991).

Ethics, data sharing, funding, and potential conflicts of interest

The study was approved by the regional ethical review board (registration number: 2020-01505). Data are available from the national Swedish spine register (Swespine) after approval by a Swedish regional ethical review board and approval by the Swespine board. There was no external source of funding for this study. The authors declare no conflicts of interest.

Results

Baseline (Table 1)

There were no statistically significant differences in BMI and mean duration of follow-up but there were some minor differences in age and sex distributions.

Reoperation rates (Figure 1, Table 2)

For spinal stenosis with DS, the reoperation rate at the index level (L4–5) was 3.0% for decompression and fusion and 6.0% for decompression only. At the 1st cranial adjacent level (L3–4), the corresponding numbers were 9.7% and 4.2% respectively. For spinal stenosis without DS, the reoperation rate at the index level (L4–5) was 3.7% for decompression and fusion and 6.2% after decompression only. At the 1st cranial adjacent level (L3–4), the corresponding numbers were 8.1% and 3.8% respectively. There were in total 38 reoperations because of implant related problems (loosening 4, breakage 1, pain because of implant 24, and pseudarthrosis 9).

Risk factors for additional surgery (Table 3)

There was a minor age difference but no statistically significant difference in BMI, sex, or fusion method for patients with DS who underwent additional surgery at the adjacent level after decompression and fusion compared with the patients that required no additional surgery at the adjacent level.
The 1-year and 2-year response rates for PROMs were 73% and 65% respectively. For all PROMs, there was no statistically significant difference between decompression and fusion and decompression only.

**Discussion**

Adjacent segment disease after lumbar spinal fusion surgery has been a topic of concern for many years. The randomized controlled trials (RCTs) of Försth et al. (2016) and Ghogawala et al. (2016) did little to settle the controversy as the results were divergent. We report reoperation rates at adjacent segments for the largest cohort of patients who underwent surgery for lumbar L4–5 spinal stenosis.

We found a statistically significant lower risk for reoperation at the index level in the fusion group compared with the decompression group. When the reoperations at index level were counted together with the reoperations at the adjacent level there were no statistically significant differences in reoperation rates between decompression and fusion and decompression only. Also, Försth et al. (2016) and Ghogawala et al. (2016) found that reoperations at the index level were more common after decompression only while reoperations at adjacent levels were more common after fusion. Our results confirm the results of Radcliff et al. (2013) that lumbar fusion is not associated with increased reoperation rate compared with decompression only.

---

**Patient-reported outcome measures (Figures 2 and 3)**

The 1-year and 2-year response rates for PROMs were 73% and 65% respectively. For all PROMs, there was no statistically significant difference between decompression and fusion and decompression only.

---

**Table 2. Results by procedure after L4–5 decompression and fusion or decompression only at the index level (L4–5), 1st cranial adjacent level (L3–4), 2nd cranial adjacent level (L2–3), and 1st caudal adjacent level (L5–S1)**

| Stenosis with DS | Reoperations, n (%) | Decompression and fusion | Decompression only |
|------------------|---------------------|--------------------------|--------------------|
|                   | n = 1,338           | n = 597                  | RR (95% CI)        |
| Index level       |                     |                          |                    |
| Excluding implant failure | 11 (0.8)  | 36 (6.0)  | 0.1 (0.07–0.3)   |
| Including implant failure  | 40 (3.0)  | 36 (6.0)  | 0.5 (0.3–0.8)    |
| Adjacent level     |                     |                          |                    |
| 1st cranial (L3–4) | 130 (9.7)  | 25 (4.2)   | 2.3 (1.5–3.5)    |
| 2nd cranial (L2–3) | 39 (2.9)   | 10 (1.7)   | 1.7 (0.9–3.5)    |
| 1st caudal (L5–S1) | 7 (0.5)    | 9 (1.5)    | 0.3 (0.1–0.9)    |
| Total              | 216 (16.1) | 80 (13.4)  | 1.2 (0.9–1.5)    |

---

**Table 3. Characteristics of patients with degenerative spondylolisthesis who underwent additional surgery at the adjacent level after L4–5 decompression and fusion compared with patients who required no additional surgery at the adjacent level**

| Parameter                      | n = 130 | n = 1,208 | p-value |
|--------------------------------|---------|-----------|---------|
| Mean age (SD)                  | 63 (8.9)| 65 (9.5)  | < 0.01  |
| BMI (SD)                       | 27 (4.0)| 27 (4.6)  | 0.8     |
| Women, n (%)                   | 103 (80)| 924 (76.5)| 0.5     |
| Fusion method, n (%)           | 103 (80)| 924 (76.5)| 0.2     |
| PLF                            | 112 (86)| 1101 (91.1)|       |
| PLIF                           | 8 (6)   | 47 (3.9)  |         |
| TLIF                           | 10 (8)  | 60 (5.0)  |         |
The overall reoperation rate reported by Sears et al. (2011) and Radcliff et al. (2013) were similar to ours. The RCT of Försth et al. (2016), however, reported higher reoperation rates but that study included multilevel procedures. Sears et al. (2011) found a correlation between the number of levels fused and reoperation rate, which might explain our somewhat lower reoperation rate. Moreover, we excluded 52 acute reoperations, which reduce our reoperation rate. Our lower reoperation rate may also be related to coverage issues of register data.

Our data suggests that the times to reoperation were shorter for reoperations at the index level compared with reoperations at the adjacent levels (Figure 1). A possible explanation is that reoperations at the index level may have several causes, e.g., insufficient decompression, instability, or implant failure, that require more urgent intervention while ASD is a more slowly progressing condition.

The reoperation rate at the 1st caudal level (L5–S1) was low and we agree with Maragkos et al. (2020) that surgeons should refrain from prophylactic procedures at the L5–S1 level when considering a posterior L4–L5 fusion.

For spinal stenosis with DS, the reoperation rate at the 1st adjacent level was 4% for decompression only (Table 2). We agree with Bydon et al. (2016) that this implies that increased rigidity across previously mobile segments introduced by spinal fusion is not the only contributing factor in the development of ASD.

The improvements in PROMs after revision surgery for ASD were similar to the minimal important change values reported by Parai et al. (2020) (Figure 3). Also Park et al. (2004) reported that surgery for ASD had relatively modest outcomes. This is an important finding in the context of patient information and shared decision-making. Patients should be informed that the results of a reoperation might be inferior in comparison with the index operation.

We found no statistically significant difference in sex or BMI for patients who required additional surgery at the adjacent level compared with the patients who required no additional surgery. Several authors have reported that sex is not associated with the development of ASD (Lee et al. 2014, Heo et al. 2015, Ou et al. 2015). Contrary to our findings, Ou et al. (2015) found an association between BMI and ASD development. That study included only 13 cases of ASD and used MRI and clinical findings (i.e., not reoperations) to define ASD.

Sears et al. (2011), Lee et al. (2014) and Heo et al. (2015) found that increasing age was a risk factor for ASD requiring further surgery. In contrast, Radcliff et al. (2013) found no association between age and reoperation rate. Although statistically significant, the difference in mean age between the groups of our study was small. If age is an important factor in
the development of ASD, it might be more correct to assess biological age or frailty rather than chronological age. Furthermore, a possible confounding factor when studying age and surgery is that younger patients may be more likely to be selected for additional surgery than older patients. This might be one part of the explanation why the patients in the reoperation group of our study are younger than the patients who did not require additional surgery.

We could not find any statistically significant association between type of fusion and additional operations at the adjacent level. In contrast, Lee et al. (2014) and Heo et al. (2015) found associations between fusion types and development of ASD, but their findings were inconsistent. Lee et al. (2014) found that interbody fusion showed a higher incidence of ASD requiring surgery than posterolateral fusion while Heo et al. (2015) reported that posterolateral fusion showed a lower survival rate compared with interbody fusion. The importance of type of surgery on the development of ASD remains to be established.

Our study has several limitations. We recognize the limitations of a register study, e.g., different implants, surgical techniques, postoperative regimen, and selection bias. In this register study, no radiographs were available for analysis. Furthermore, higher response rates for the PROMs would have strengthened our results but studies following non-responders have revealed similar results in terms of the PROMs (Solberg et al. 2011). Our primary outcome variable, the reoperation rate, was of course not affected by the missing PROMs. Moreover, to present our data in a simple and transparent way we did not adjust the reoperation rates and the PROMs for differences in baseline data by using advanced statistical methods, e.g., regression analysis or propensity score matching. Given the diverging results of previous studies (Sears et al. 2011, Radcliff et al. 2013) concerning age as a confounding factor in relation to ASD, we find unlikely that the minor age differences in our groups have any substantial impact on the results.

In conclusion, single-level lumbar fusion surgery at the L4–5 level is associated with an increased rate of reoperations at the proximal adjacent level compared with decompression only. However, when reoperations on the index level are included there is no difference in reoperation rates between fusion and decompression only. Addressing the index segment for durable outcome should be the primary objective of surgery. This can be achieved with decompression or decompression and fusion with similar risk for additional surgery in selected cases.

Study design: AJ, FGS. Analysis of data: AJ. Interpretation of data: AJ, FN, MH, FGS. Drafting the manuscript: AJ, FGS. Critically revising the manuscript: FN, MH, FGS.

Acta Orthopaedica 2021; 92 (3): 264–268

Altmann D G. Practical statistics for medical research. London: Chapman and Hall; 1991.

Bydon M, Macki M, De la Garza-Ramos R, McGovern K, Sciubba D M, Wolinsky J P, Witham T F, Gokaslan Z L, Bydon A. Incidence of adjacent segment disease requiring reoperation after lumbar laminectomy without fusion: a study of 398 patients. Neurosurgery 2016; 78(2): 192–9.

Dolan P. Modeling valuations for EuroQol health states. Med Care 1997; 35(11): 1095–108.

EuroQol Group. EuroQol: a new facility for the measurement of health-related quality of life. Health Policy 1990; 16(3): 199-208.

Fairbank J C, Pynsent P B. The Oswestry disability index. Spine (Phila Pa 1976) 2000; 25(22): 2940-52.

Förgh P, Olafsson G, Carlsson T, Frost A, Borgström F, Fritzell P, Ohagen P, Michaelsson K, Sandén B. A randomized, controlled trial of fusion surgery for lumbar spinal stenosis. N Engl J Med 2016; 374(15): 1413-23.

Ghogawala Z, Dziura J, Butler W E, Dai F, Terrin N, Magge S N, Coumans J V C E, Harrington J F, Amin-Hanjani S, Schwartz J S, Sonntag V K H, Barker F G 2nd, Benzell E C. Laminctomy plus fusion versus laminectomy alone for lumbar spondylolisthesis. N Engl J Med 2016; 374(15): 1424-34.

Ghogawala Z, Resnick D K, Glassman S D, Dziura J, Shaffrey C I, Mummaneni P V. Randomized controlled trials for degenerative lumbar spondylolisthesis: which patients benefit from lumbar fusion? J Neurosurg Spine 2017; 26(2): 260-6.

Heo Y, Park J H, Seong H Y, Lee Y S, Jeon S R, Rhim S C, Roh S W. Symptomatic adjacent segment degeneration at the L3–4 level after fusion surgery at the L4–5 level: evaluation of the risk factors and 10-year incidence. Eur Spine J 2015; 24(11): 2474-80.

Lee J C, Kim Y, Soh J W, Shin B J. Risk factors of adjacent segment disease requiring surgery after lumbar spinal fusion: comparison of posterior lumbar interbody fusion and posterolateral fusion. Spine (Phila Pa 1976) 2014; 39(5): E339-45.

Maragkos G A, Atesok K, Papavasiliou E. Prognostic factors for adjacent segment disease after L4-L5 lumbar fusion. Neurosurgery 2020; 86(6): 835-42.

Okuda S, Nagamoto Y, Matsumoto T, Sugiuira T, Takahashi Y, Iwasaki M. Adjacent segment disease after single segment posterior lumbar interbody fusion for degenerative spondylolisthesis: minimum 10 years follow-up. Spine (Phila Pa 1976) 2018; 43(23): E1384-E1388.

Ou C Y, Lee T C, Lee T H, Huang Y H. Impact of body mass index on adjacent segment disease after lumbar fusion for degenerative spine disease. Neurosurgery 2015; 76(4): 396-401.

Parai C, Hägg O, Lind B, Brisby H. ISSLS prize in clinical science 2020: The reliability and interpretability of score change in lumbar spine research. Eur Spine J 2020; 29(4): 663-9.

Park P, Barton H J, Gala V C, Hoff J T, McGillicuddy J E. Adjacent segment disease after lumbar or lumbosacral fusion: Review of the literature. Spine (Phila Pa 1976) 2004; 29(17): 1938-44.

Radcliff K, Curry P, Hilibrand A, Kepler C, Lurie J, Zhao W, Albert T J, Weinstein J. Risk for adjacent segment and same segment reoperation after surgery for lumbar stenosis: a subgroup analysis of the Spine Patient Outcomes Research Trial (SPORT). Spine (Phila Pa 1976) 2013; 38(7): 531-9.

Sears W R, Sergièdes I G, Kazemi N, Smith M, White G J, Osburg B. Incidence and prevalence of surgery at segments adjacent to a previous posterior lumbar arthrodesis. Spine 2011; 36(11): 11-20.

Solberg T K, Mendeloff A, Vaishnav A S, Othman Y, Virk S, Katsumura Y, Iyer S, McNaney S, Albert T, Gang C H, Qureshi S A. The necessity and risk factors of subsequent fusion after decompression alone for lumbar spinal stenosis with lumbar spondylolisthesis: 5 years follow-up in 2 different large populations. Spine J 2020; 20(10): 1566-72.