

Four-year follow-up of surgical versus non-surgical therapy for chronic low back pain

Jens Ivar Brox,¹ Øystein P Nygaard,² Inger Holm,³ Anne Keller,⁴ Tor Ingebrigtsen,⁵ Olav Reikerås⁶

► Additional data are published online only. To view these files please visit the journal online (<http://ard.bmj.com>).

¹Orthopaedic Department, Oslo University Hospital – Rikshospitalet, Oslo, Norway

²Neurosurgical Department, St Olavs Hospital, Trondheim, Norway

³Clinic of Rehabilitation, Oslo University Hospital – Rikshospitalet, Oslo, Norway

⁴Department of Physical Medicine and Rehabilitation, Oslo University Hospital – Ullevaal, Oslo, Norway

⁵Neurosurgical Department, University Hospital of North Norway, Tromsø, Norway

⁶Hospital of Rehabilitation, Stavern and Unifob Health, University of Bergen, Bergen, Norway

Correspondence to

Dr Jens Ivar Brox, Orthopaedic Department, Oslo University Hospital-Rikshospitalet, Sognsvannsveien, 0027 Oslo, Norway; jens.ivar.brox@rikshospitalet.no

Accepted 26 June 2009

ABSTRACT

Objectives To compare the long-term effectiveness of surgical and non-surgical treatment in patients with chronic low back pain.

Methods Two merged randomised clinical trials compared instrumented transpedicular fusion with cognitive intervention and exercises in 124 patients with disc degeneration and at least 1 year of symptoms after or without previous surgery for disc herniation. The main outcome measure was the Oswestry disability index.

Results At 4 years 14 (24%) patients randomly assigned to cognitive intervention and exercises had also undergone surgery. 15 (23%) patients assigned fusion had undergone re-surgery. The mean treatment effect for the primary outcome was 1.1; 95% CI –5.9 to 8.2, according to the intention-to-treat analysis and –1.6; 95% CI –8.9 to 5.6 in the as-treated analysis. There was no difference in return to work.

Conclusions Long-term improvement was not better after instrumented transpedicular fusion compared with cognitive intervention and exercises.

Lumbar spine fusion for chronic low back pain has increased rapidly during the past two decades.¹ Four randomised studies have compared lumbar fusion and conservative treatment in patients with disc degeneration and chronic low back pain.^{2–5} Results up to 2 years after treatment have been published. A recent meta-analysis concluded that cumulative evidence at the present time does not support routine fusion, whereas a recent systematic review concluded that surgery may be more efficacious than unstructured care, but may not be more efficacious than structured cognitive-behavioural therapy.^{6,7} Methodological limitations of the randomised trials prevent firm conclusions. The Norwegian studies, published in three papers,^{4,5,8} were criticised for lack of power, short follow-up and a high number of withdrawals from fusion among patients with chronic low back pain after surgery for disc herniation. Because results, interventions and outcome measures were similar we merged the two Norwegian trials for long-term follow-up using a questionnaire mailed to the patients. We report the 4-year effectiveness of lumbar fusion compared with cognitive intervention and exercises in patients with chronic low back pain with and without previous surgery for disc herniation.

PATIENTS AND METHODS

Study design

The Norwegian studies were investigator initiated in 1999 and were conducted at four university hospitals. They were designed as two separate

randomised trials and results were reported after 1-year follow-up.^{4,5} The ethics committee for medical research in health region I of Norway approved the studies.

Patients

Patients aged 25–60 years with chronic low back pain for at least 1 year, Oswestry disability index score greater than 30, and disc degeneration at L4–L5 and/or L5–S1, were eligible to participate in the study. Exclusion criteria were: widespread myofascial pain; spinal stenosis with reduced walking distance and neurological signs; disc herniation or lateral recess stenosis with clinical signs of radiculopathy; inflammatory disease; previous spinal fracture; previous fusion surgery of the spine; pelvic pain; generalised disc degeneration on plain radiographic examination; ongoing serious somatic and psychiatric disease; registered medicine abuse and reluctance to accept one of the interventions. At least one spine surgeon and one specialist in physical medicine and rehabilitation examined each patient. A research physiotherapist coordinated the study and verified eligibility. All eligible patients were given oral and written information about the study and the two interventions.

Randomisation

Patients received treatment assignments from an independent unit at Unifob Health, University of Bergen that was not involved in the treatment. Computer-generated randomly permuted blocks were used and allocation was concealed. The project coordinator telephoned the unit at Unifob Health and reported an identification number and was phoned back in order to inform the patient about the assigned intervention.⁴ Treatments were started within 3 months after randomisation.

Study interventions

The protocol surgery was posterolateral fusion with transpedicular screws of the L4–L5 and/or L5–S1 segment. Autologous bone was used in all cases. Postoperative rehabilitation was at the choice of the surgeon. Surgery was performed at two neurosurgical and two orthopaedic departments.⁴

The cognitive intervention and exercises consisted of 1 week plus 2 weeks in the outpatient clinic at the study centre interrupted by 2 weeks at home. Specialists in physical medicine and physiotherapists gave the intervention. In addition, patients met a peer for exchanging experiences. The main aim was to make the patients confident



This paper is freely available online under the BMJ Journals unlocked scheme, see <http://ard.bmj.com/info/unlocked.dtl>

that they could not do any harm to the disc (back) by engaging in ordinary activities of daily life. Details of the programme have been outlined previously.⁴

Outcome measures

A standardised questionnaire was sent by post to all patients. The primary outcome measure was the original (version 1.0) Oswestry disability index.⁹ This score has 10 questions about pain and disability and ranges from 0% (no pain and disability) to 100% (worst possible disability).

Secondary outcome measures included pain,³ general function score,¹⁰ global back disability question for the assessment of patients' overall rating,¹¹ work and medication,¹¹ emotional distress,¹² fear-avoidance beliefs¹³ and life satisfaction (for details see additional supplemental file, available online only).¹⁴ The questionnaire also included questions about treatment taken after the 1-year follow-up. Additional surgery was verified from medical records.

Statistical analysis

Estimation of sample sizes in the two trials merged for 4-year follow-up has been reported previously.^{4,5} Results are primarily

analysed with an intention-to-treat approach. Because of cross-over and withdrawal, sensitivity analyses were based on the treatment actually received. Baseline characteristics in those who attended the 4-year follow-up were compared with cross-over patients and withdrawals in the two treatment groups (table 1). Means (\pm SD) or numbers (percentages) were calculated for baseline and 4-year follow-up in those who attended, and are reported separately for intention-to-treat and as-treated analyses. The analyses of treatment effects compared differences between interventions at 4 years using linear regression with adjustments for gender, age, previous surgery for disc herniation and baseline scores. We conducted analyses with and without the most recent observed non-missing value carried forward in those who did not attend the 4-year follow-up. We used this simplistic method, being aware that more comprehensive multiple imputation techniques are available.¹⁵ The estimated treatment effects are reported as mean adjusted differences between groups (95% CI) based on analyses using the last observed value carried forward and including all patients randomly assigned (tables 2 and 3). Categorical outcomes (patients' overall rating, medication and work) were dichotomised and logistic regression was used to calculate adjusted OR (95% CI) with adjustments for gender, age, previous surgery for disc herniation and

Table 1 Baseline characteristics of the patients*

	Lumbar fusion		Cognitive intervention and exercises	
	All randomised (n=66)	Crossover/withdrawals (n=11)†	All randomised (n=58)	Crossover/withdrawals (n=17)†
Age (years)	42.7 \pm 8.0	43.9 \pm 7.3	42.4 \pm 8.0	42.1 \pm 7.7
No of men (%)	27 (41)	7 (64)	29 (50)	8 (47)
Years from first pain episode	8.9 \pm 7.9	8.1 \pm 7.9	9.6 \pm 7.4	12.2 \pm 9.6
Married/living together no (%)	57 (86)	10 (91)	49 (81)	15 (88)
Occupational education <3 years no (%)	45 (68)	8 (27)	38 (66)	9 (53)
Work status no (%)				
Working	9 (14)	2 (18)	9 (16)	1 (6)
On sick leave	14 (21)	3 (27)	16 (28)	5 (29)
On rehabilitation	29 (44)	3 (27)	22 (38)	7 (41)
Disability pension	10 (15)	3 (27)	10 (17)	4 (24)
Student, homemaker, unemployed	3 (5)		1 (2)	
Retirement pension	1 (2)			
Back pain (0–100)‡	63.0 \pm 14.7	64.2 \pm 15.5	64.6 \pm 12.5	65.2 \pm 12.0
Oswestry disability index§	44.5 \pm 10.7	45.3 \pm 10.1	44.2 \pm 11.0	47.0 \pm 7.8
Emotional distress (1–4)¶	1.9 \pm 0.5	1.9 \pm 0.7	1.9 \pm 0.5	1.9 \pm 0.5
Previous surgery for disc herniation no (%)	29 (44)	4 (36)	31 (53)	8 (47)
Beliefs in surgery**	69.7 \pm 18.2	62.1 \pm 17.7	72.4 \pm 20.3	70.3 \pm 16.5
Beliefs in non-surgical treatment**	40.1 \pm 25.4	42.1 \pm 24.5	44.5 \pm 25.1	44.0 \pm 24.7
Comorbidity no (%)	24 (36)	6 (55)	18 (31)	6 (35)
Taking analgesics daily or weekly no (%)	40 (61)	4 (36)	40 (69)	16 (94)
Smoking no (%)	36 (55)	4 (36)	30 (52)	9 (53)

*Plus–minus values are means \pm SD.

†Three patients allocated lumbar fusion died, four did not have and four had cognitive intervention and exercises. Fourteen patients allocated cognitive intervention and exercises had surgery.

‡Back pain ranges from 0 to 100, with lower scores indicating less severe symptoms.

§The Oswestry disability index ranges from 0 to 100, with lower scores indicating less severe symptoms.

¶Emotional stress ranges from 1 to 4, with lower scores indicating less severe symptoms.

**Beliefs ranges from 0 to 100, with lower scores indicating not efficient.

baseline scores.⁴ Analyses were performed with the use of SPSS software, version 15.

RESULTS

Patients

A total of 124 patients was enrolled out of 234 who were eligible: 66 were assigned to the surgical group and 58 to the non-surgical group (figure 1). The 4-year follow-up rate was 92% and 86%, respectively. In the surgical group, 88% had undergone surgery at 1 year and 91% at 4 years. In the non-surgical group, 5% had undergone surgery at 1 year and 24% at 4 years.

In both groups patients had stronger beliefs in surgical compared with non-surgical treatment at baseline (table 1). Crossover patients and withdrawals from surgery were more

often men and non-smokers, had higher occupational education and higher comorbidity, but took analgesics less often at baseline. Such patients from the non-surgical group took analgesics more often at baseline.

Healthcare utilisation and return to work

Thirty (49%) and 29 (58%) allocated surgical or non-surgical treatment, respectively, reported visits to a physician for back pain the year before the 4-year follow-up. Physiotherapy (20% vs 22%) and other treatments (16% vs 14%) were taken by a minority in both groups. More patients who had surgery (53% vs 32%) were on disability pension (adjusted OR 2.5; 95% CI 1.1 to 5.9). For the intention-to-treat analysis this difference was no longer significant ($p=0.21$). The number of patients working full time was not significantly different (tables 2 and 3).

Table 2 Intention-to-treat analysis*

Outcome	Lumbar fusion (N=61)†	Cognitive/exercises (N=50)†	Adjusted treatment effect (95% CI)‡
Primary			
Oswestry disability index†			
Baseline	44.1±10.7	43.4±11.1	1.1 (−5.9 to 8.2)
4 Years	29.7±20.5	27.0±19.4	
Secondary			
General function score†			
Baseline	37.3±19.3	40.0±18.9	−3.5 (−11.6 to 4.6)
4 Years	25.8±24.7	21.4±21.5	
Back pain†			
Baseline	62.8±14.5	64.2±12.5	2.3 (−6.4 to 10.9)
4 Years	42.2±23.9	44.7±22.8	
Lower limb pain†			
Baseline	48.5±24.4	44.8±23.5	1.3 (−8.3 to 10.8)
4 Years	34.8±29.4	33.5±24.7	
Emotional distress§			
Baseline	1.9±0.5	1.9±0.5	−0.1 (−0.1 to 0.3)
4 Years	1.7±0.6	1.7±0.6	
Life satisfaction¶			
Baseline	5.0±2.2	4.6±1.7	−0.3 (−1.1 to 0.6)
4 Years	6.2±2.5	6.4±2.3	
Fear-avoidance beliefs physical activity**			
Baseline	13.0±5.0	15.4±5.0	−3.5 (−5.8 to −1.1)
4 Years	9.1±7.3	7.0±6.0	
Fear-avoidance beliefs work**			
Baseline	26.1±10.5	28.4±10.7	−4.3 (−8.3 to −0.2)
4 Years	23.9±13.8	21.1±12.5	
Patients overall rating – no (%) success††			
1 Year	38 (62)	32 (64)	1.0 (0.8 to 1.5)
Work – no (%)			
Baseline	9 (15)	8 (16)	0.8 (0.3 to 1.9)
1 Year	16 (26)	17 (34)	

*Mean values±SD unless otherwise noted.

†The Oswestry disability index, the general function score, back and lower limb pain ranges from 0 to 100, with lower scores indicating fewer symptoms.

‡The treatment effect is the difference between patients randomly assigned to lumbar fusion and cognitive intervention and exercises at 4 years with adjustments for baseline score, age, gender and previous disc surgery. All patients randomly assigned ($n=124$) are included with last observed value carried forward.

††All who did not attend the 4-year follow-up are classified as non-success.

§Emotional stress ranges from 1 to 4, with lower scores indicating less severe symptoms.

¶Life satisfaction ranges from 1 to 10, with higher scores indicating better life satisfaction.

**Fear-avoidance beliefs for physical activity ranges from 0 to 24 and for work from 0 to 42, with lower scores indicating less strong beliefs for physical activity and work hurting the back.

Crossover, complications and re-operations

Non-adherence was registered in 17 (29%) patients randomly assigned to cognitive intervention and exercises, three (5%) did not have the allocated treatment and 14 (24%) patients later had surgery (figure 1). Eleven (17%) patients randomly assigned to surgery were classified as non-adherent, six (9%) did not have lumbar fusion (figure 1), two (3%) withdrew and three (5%) patients died. Deaths were not related to the surgical procedures. Four crossover patients operated (25%) in the non-surgical group and 15 (25%) in the surgical group had re-operation.

The reason was persistent complaints or deterioration of the condition. Complications have been described previously.^{4 5} No major complications occurred in patients operated after the 1-year follow-up.

Main treatment effects

In the intention-to-treat analysis there was no treatment effect for the Oswestry disability index. When adjusted for age, gender, baseline score and previous disc surgery the treatment effect was 1.1; 95% CI -5.9 to 8.2 (table 2). The mean adjusted treatment effect was -1.6; 95% CI -8.9 to 5.6 (table 3) according to as-treated analysis. Sensitivity analyses including only those who attended the 4-year follow-up did not alter the results.

Secondary outcome

The only treatment effect observed in the secondary outcome was a reduction of fear-avoidance beliefs favouring cognitive intervention and exercises (tables 2 and 3). The mean treatment effect for fear-avoidance beliefs for physical activity was -3.5;

Table 3 As-treated analyses*

Outcome	Lumbar fusion (N=62)	Cognitive/exercises (N=49)	Adjusted treatment effect (95% CI)†
Primary			
Oswestry disability index‡			
Baseline	44.4 ± 10.6	43.0 ± 11.1	-1.6 (-8.9 to 5.6)
4 Years	29.1 ± 20.2	27.7 ± 19.9	
Secondary			
General function score‡			
Baseline	37.9 ± 19.3	39.2 ± 19.2	-3.2 (-11.4 to 5.0)
4 Years	24.8 ± 24.7	22.5 ± 22.9	
Back pain‡			
Baseline	63.3 ± 14.7	64.2 ± 12.5	4.1 (-4.7 to 12.8)
4 Years	40.5 ± 23.0	46.8 ± 23.4	
Lower limb pain‡			
Baseline	48.4 ± 23.1	44.8 ± 24.9	-2.9 (-12.6 to 6.7)
4 Years	35.5 ± 27.7	32.7 ± 26.8	
Emotional distress§			
Baseline	1.8 ± 0.5	1.9 ± 0.5	0.1 (-0.2 to 0.3)
4 Years	1.6 ± 0.5	1.6 ± 0.7	
Life satisfaction¶			
Baseline	5.0 ± 2.3	4.7 ± 1.7	0.2 (-0.6 to 1.0)
4 Years	6.4 ± 2.4	6.2 ± 2.4	
Fear-avoidance physical activity**			
Baseline	13.1 ± 5.0	15.5 ± 5.1	-2.8 (-5.3 to -0.4)
4 Years	8.8 ± 7.0	7.3 ± 6.4	
Fear – avoidance work**			
Baseline	26.4 ± 10.1	28.1 ± 12.4	-4.8 (-8.9 to -0.7)
4 Years	23.8 ± 13.0	21.1 ± 13.4	
Patients overall rating – no (%) success††			
1 Year	38 (61)	32 (65)	1.1 (0.8 to 1.8)
Work – no (%)			
Baseline	8 (13)	9 (18)	0.8 (0.4 to 1.6)
1 Year	15 (26)	16 (33)	

*Mean values ± SD unless otherwise noted.

†The treatment effect is the difference between the lumbar fusion group and the cognitive intervention and exercises group at 4 years with adjustments for baseline score, age, gender and previous disc surgery. All patients randomly assigned (n=124) are included with last observed value carried forward.

‡The Oswestry disability index, the general function score, back and lower limb pain ranges from 0 to 100, with lower scores indicating fewer symptoms.

§Emotional stress ranges from 1 to 4, with lower scores indicating less severe symptoms.

¶Life satisfaction ranges from 1 to 10, with higher scores indicating better life satisfaction.

**Fear-avoidance beliefs for physical activity ranges from 0 to 24 and for work from 0 to 42, with lower scores indicating less strong beliefs for physical activity and work hurting the back.

††All who did not attend the 4-year follow-up are classified as non-success.

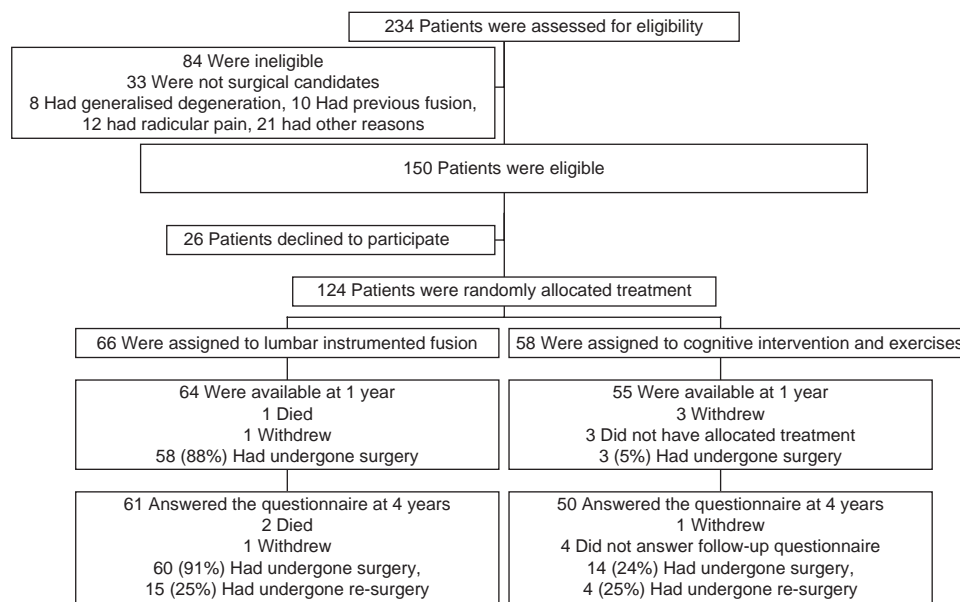


Figure 1 Exclusion, enrolment, randomisation and follow-up of participants.

95% CI -5.8 to -1.1 in the intention-to-treat analysis and -2.8 ; 95% CI -5.3 to -0.4 in the as-treated last analysis, and -4.3 ; 95% CI -8.3 to -0.2 and -4.8 ; 95% CI -8.9 to -0.7 for fear-avoidance beliefs for work, respectively. Pain medication was taken daily or weekly by 58% treated with surgery compared with 35% not operated (adjusted OR 2.3; 95% CI 1.0 to 5.2). For the intention-to-treat analysis the difference was no longer significant ($p=0.14$).

DISCUSSION

In patients with chronic low back pain with and without previous surgery for disc herniation, lumbar fusion was not superior to cognitive intervention and exercises at relieving symptoms, improving function and return to work at 4-years. The results were consistent for intention-to-treat and as-treated analyses. The number of re-operations in patients randomly assigned to surgery were similar to the number patients operated in the non-surgical group.

The CI for the treatment effects were within 10 points on the Oswestry disability index that the trial was designed to detect. This indicates that lack of power is unlikely to explain the observed results.

Comparison with existing literature

The present study is the first to provide the long-term results of a randomised study comparing lumbar fusion with non-surgical treatment in patients with chronic low back pain. Results are in agreement with previously reported results at 1 and 2 years.²⁻⁵ The reported long-term results do not exclude the possibility that fusion may be indicated in carefully selected patients with chronic low back pain, but widening indications have contributed to the rise in rates of fusion surgery.¹⁶ Despite much effort to improve selection criteria, there is no agreement on providing a valid tool to diagnose discogenic pain, and even procedures such as discography and MRI are not reliable for selecting patients.¹⁷ Hägg *et al*¹⁸ reported that a personality characterised by low neuroticism and low disc height predicted functional improvement after surgery and that work resumption was predicted by low age and short-term sick leave.

The re-operation rate was slightly higher than previously reported after spinal surgery.¹⁹ A higher rate is not unexpected after instrumented fusion compared with laminectomy and discectomy. Although device failure and postoperative infection did not explain the cases in the present study, the outcome after surgery for back pain may be less predictable than after surgery for leg pain. Re-operation is an undesirable outcome, and the high rate observed in the present study is an argument against surgery. Preventing repeat spinal surgery is an important goal for surgeons and patients.

We observed no treatment effects in secondary outcome except for fear-avoidance beliefs. Differences in favour of non-surgical treatment for the number taking pain medication regularly or on disability pension were observed in the as-treated analysis only. The aim of the non-surgical intervention was to give patients the understanding that they could not do any harm to the disc (back) by engaging in ordinary activities of daily life. To reduce fear and avoidance and achieve confidence patients were encouraged and confronted with physical activities that were previously not recommended. Results at 4 years suggest that the reduction in avoidant behaviour observed at 1 year was maintained.

We observed that more patients used pain medication after surgery. Alternative interpretations are that these patients either experience more pain or they are habituated to pain medication. A recent study reported that surgical patients used more opiates, but that both pain medication and pain intensity were reduced after participation in a multidisciplinary pain programme.²⁰ The reduction was attributed to a cognitive-behavioural approach to symptom management during the course of the rehabilitation programme. The possible effect of multidisciplinary pain rehabilitation on withdrawal of pain medication warrants further studies.

Most patients included were out of work at baseline. The number who had returned to work was not significantly different at 4 years, but the OR for disability pension was increased after surgery. Our interpretation is that the claim adjuster may consider that lumbar fusion represents the end stage of treatment, and consequently the claim for disability pension may be more easily accepted.

Possible confounders and weaknesses

A limitation of this study is the non-adherence to randomised treatment. Although patients consented to the protocol, some of them chose to change their consent as they are allowed to in clinical trials. The degree of non-adherence was lower than in the SPORT studies.^{21 22} One possible interpretation is that we aimed to conduct the interventions within 3 months after enrolment compared with 3–6 months in the SPORT studies. The consistency of results in intention-to-treat and as-treated analyses of the present study indicates that non-adherence does not play a decisive role to explain our results. Although 89% answered the follow-up questionnaire, the use of last value carried forward and not the multiple imputation technique for missing values may bias results.

Another limitation is the lack of a placebo group. Expectations are important for outcome. Sham surgery has previously shown that methods expected to be highly effective were mediated by placebo.²³ We are unable to exclude the possibility that the observed improvements reflect the natural course, placebo or expectations and care.

Surgeons, patients and stakeholders may consider new technical surgical solutions more powerful, implying fast improvement and simple technical solutions in the hands of a skilled spinal surgeon, but postulated advantages for new procedures are based more on theories than knowledge.²⁴ The introduction of new technology in clinical practice should be based on sound evidence from randomised studies.²⁵ Patients allocated non-surgical treatment should be given the best evidence intervention and the same attention and care as the surgical patients.

In conclusion, patients did not have a better long-term improvement after instrumented fusion compared with cognitive intervention and exercises.

Acknowledgements The authors would like to thank the patients who participated in the trial, the nurses and the nurse aids at the hospital departments and outpatient clinics and the referring medical doctors: A Friis for coordinating inclusion, treatments and 1-year follow-up; H Ursin and H Eriksen at Unifob Health, University of Bergen for their work with the random assignment of patients and comments on study design; AH Pripp at Rikshospitalet University Hospital for statistical advice; physiotherapists AK Koller, MFosdahl and T Haakenstad for non-surgical treatments; R Sørensen, JE Lange, R Riise and O Grundnes for lumbar fusions and the radiologists R Gunderson and AM Finnanger for their assistance.

Competing interests None.

Patient consent Obtained.

Ethics approval This study was conducted with the approval of the Ethics Committee Health region I, Norway.

Provenance and peer review Not commissioned; externally peer reviewed.

REFERENCES

1. Cowan JA Jr, Dimick JB, Wainess R, *et al*. Changes in the utilization of spinal fusion in the United States. *Neurosurgery* 2006;**59**:15–20; discussion 15–20.
2. Fairbank J, Frost H, Wilson-MacDonald J, *et al*. Randomised controlled trial to compare surgical stabilisation of the lumbar spine with an intensive rehabilitation programme for patients with chronic low back pain: the MRC spine stabilisation trial. *BMJ* 2005;**330**:1233–9.
3. Fritzell P, Hägg O, Wessberg P, *et al*. 2001 Volvo Award Winner in Clinical Studies: Lumbar fusion versus nonsurgical treatment for chronic low back pain: a multicenter randomized controlled trial from the Swedish Lumbar Spine Study Group. *Spine* 2001;**26**:2521–32; discussion 2532–4.
4. Brox JI, Sørensen R, Friis A, *et al*. Randomized clinical trial of lumbar instrumented fusion and cognitive intervention and exercises in patients with chronic low back pain and disc degeneration. *Spine* 2003;**28**:1913–21.
5. Brox JI, Reikerås O, Nygaard Ø, *et al*. Lumbar instrumented fusion compared with cognitive intervention and exercises in patients with chronic back pain after previous surgery for disc herniation: a prospective randomized controlled study. *Pain* 2006;**122**:145–55.
6. Ibrahim T, Tleyjeh IM, Gabbar O. Surgical versus non-surgical treatment of chronic low back pain: a meta-analysis of randomised trials. *Int Orthop* 2008;**32**:107–13.
7. Mirza SK, Deyo RA. Systematic review of randomized trials comparing lumbar fusion surgery to nonoperative care for treatment of chronic back pain. *Spine* 2007;**32**:816–23.
8. Keller A, Brox JI, Gunderson R, *et al*. Trunk muscle strength, cross-sectional area, and density in patients with chronic low back pain randomized to lumbar fusion or cognitive intervention and exercises. *Spine* 2004;**29**:3–8.
9. Fairbank JC, Couper J, Davies JB, *et al*. The Oswestry low back pain disability questionnaire. *Physiotherapy* 1980;**66**:271–3.
10. Hägg O, Fritzell P, Romberg K, *et al*. The General Function Score: a useful tool for measurement of physical disability. Validity and reliability. *Eur Spine J* 2001;**10**:203–10.
11. Holm I, Friis A, Storheim K, *et al*. Measuring self-reported functional status and pain in patients with chronic low back pain by postal questionnaires: a reliability study. *Spine* 2003;**28**:828–33.
12. Derogatis LR, Lipman RS, Rickels K, *et al*. The Hopkins Symptom Checklist (HSCL): a self-report symptom inventory. *Behav Sci* 1974;**19**:1–15.
13. Waddell G, Newton M, Henderson I, *et al*. A Fear-Avoidance Beliefs Questionnaire (FABQ) and the role of fear-avoidance beliefs in chronic low back pain and disability. *Pain* 1993;**52**:157–68.
14. Andrews FM, Robinson JP. *Measures of subjective well-being*. In: Robinson JP, Shaver PR, Wrightsman LS, eds. *Measures of personality and social psychological attitudes*. San Diego, California, USA: Academic Press, 1991.
15. Carpenter JR, Kenward MG, Vansteelandt S. A comparison of multiple imputation and doubly robust estimation for analysis with missing data. *J Roy Statist Soc (A)* 2006;**169**:1–14.
16. Deyo RA, Nachemson A, Mirza SK. Spinal-fusion surgery – the case for restraint. *N Engl J Med* 2004;**350**:722–6.
17. Carragee EJ. Clinical practice. Persistent low back pain. *N Engl J Med* 2005;**352**:1891–8.
18. Hägg O, Fritzell P, Ekselius L, *et al*. Predictors of outcome in fusion surgery for chronic low back pain. A report from the Swedish Lumbar Spine Study. *Eur Spine J* 2003;**12**:22–33.
19. Martin BI, Mirza SK, Comstock BA, *et al*. Are lumbar spine reoperation rates falling with greater use of fusion surgery and new surgical technology? *Spine* 2007;**32**:2119–26.
20. Crisostomo RA, Schmidt JE, Hooten WM, *et al*. Withdrawal of analgesic medication for chronic low-back pain patients: improvement in outcomes of multidisciplinary rehabilitation regardless of surgical history. *Am J Phys Med Rehabil* 2008;**87**:527–36.
21. Weinstein JN, Tosteson TD, Lurie JD, *et al*. Surgical versus nonsurgical therapy for lumbar spinal stenosis. *N Engl J Med* 2008;**358**:794–810.
22. Weinstein JN, Lurie JD, Tosteson TD, *et al*. Surgical versus nonsurgical treatment for lumbar degenerative spondylolisthesis. *N Engl J Med* 2007;**356**:2257–70.
23. Chalmers TC. Randomization and coronary artery surgery. *Ann Thorac Surg* 1972;**14**:323–7.
24. Gibson JN, Waddell G. Surgery for degenerative lumbar spondylosis. *Cochrane Database Syst Rev* 2005;**4**:CD001352.
25. Weinstein JN. The tortoise and the hare: is there a place in spine surgery for randomized trials? *Spine* 1999;**24**:2548–9.