Surgery versus prolonged conservative treatment for sciatica: 5-year results of a randomised controlled trial

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ABSTRACT

Objective: This study describes the 5 years’ results of the Sciatica trial focused on pain, disability, (un)satisfactory recovery and predictors for unsatisfactory recovery.

Design: A randomised controlled trial.

Setting: Nine Dutch hospitals.

Participants: Five years’ follow-up data from 231 of 283 patients (82%) were collected.

Intervention: Early surgery or an intended 6 months of conservative treatment.

Main outcome measures: Scores from Roland disability questionnaire, visual analogue scale (VAS) for leg and back pain and a Likert self-rating scale of global perceived recovery were analysed.

Results: There were no significant differences between groups on the 5 years’ primary outcome scores. Despite at least 6 months of conservative treatment 46% of the conservatively allocated patients were treated surgically because of severe leg pain and disability. Forty-nine (21%) patients had an unsatisfactory recovery at 5 years and the recovery pattern showed that there was a variable group of 66 patients (31%) with at least one unsatisfactory outcome at 1, 2 or 5 years of follow-up. Multivariate logistic regression showed that age (>40; OR 2.42 (95% CI 1.16 to 5.02)), severity of leg pain (VAS >70; OR 3.32 (95% CI 1.69 to 6.54)) and the Mc Gill affective score (score >3; OR 6.23 (95% CI 2.23 to 17.38)) were the only significant predictors for an unsatisfactory outcome at 5 years.

Conclusions: In the long term, 8% of the patients with sciatica never showed any recovery and in at least 23%, sciatica appears to result in ongoing complaints, which fluctuate over time, irrespective of treatment. Prolonged conservative care might give patients a fair chance for pain and disability to resolve without surgery, but with the risk to receive delayed surgery after prolonged suffering of sciatica. Age above 40 years, severe leg pain at baseline and a higher affective McGill pain score were predictors for unsatisfactory recovery. Trial Registry ISRCT No 26872154.

INTRODUCTION

The lumbosacral radicular syndrome (LSRS), caused by a herniated lumbar disc, is one of the most expensive disorders for society in terms of work absenteeism and disability.1 In 2007, the Sciatica Trial showed that the clinical outcome after 1 year was not different...
from prolonged conservative treatment, although recovery within the first year was better with early surgery. In the prolonged conservative treatment group, however, 39% of patients crossed over to surgical treatment because of intractable pain within 1 year and 44% within 2 years of follow-up. Despite the fact that this study showed, along with other randomised controlled trials, that a strategy of prolonged conservative care is safe and reduces the risk for patients undergoing surgery, the optimal timing of surgery with regard to long-term outcome has still not been defined.

Although LSRS is described in the literature as having a quite favourable course, one might question this assumption as the 2 years’ follow-up showed that about 20% of patients report an unsatisfactory outcome on all outcome scales and that the risk to suffer prolonged disability is higher than expected beforehand.

The primary aim of the present study was to compare the pain and disability scores at 5 years’ follow-up between patients in the Sciatica Trial randomised for surgery or randomised for prolonged conservative treatment. The second aim was to evaluate the proportion of patients with an unsatisfactory recovery at 5 years’ follow-up and to identify factors contributing to these unsatisfactory results.

**MATERIAL AND METHODS**

**Design**

The study is part of the Sciatica Trial, a multicentre, prospective randomised trial among patients with 6–12 weeks of sciatica to determine whether a strategy of early surgery leads to better outcomes during the first year than does a strategy of prolonged conservative treatment for an additional 6 months followed by surgery for those patients who do not improve.

In summary, patients, 18–65 years of age, with an LSRS with a concomitant disc herniation confirmed by MRI, were eligible for participation. A computer-generated permuted-block scheme was used for randomisation stratified by centre. In the surgical intervention group, a disc herniation removal through a unilateral transflaval approach using optical magnification was performed. Prolonged conservative treatment regimen was defined by general practitioners and treatment was mainly aimed at resuming daily activities. Patients were notified beforehand that they were participating in a study comparing two different strategies for the timing of intervention rather than comparing surgery with non-surgical treatment.

The design and study protocol have been published previously. Baseline characteristics from the Sciatica Trial have been published previously and were combined with the findings obtained from the 5 years’ follow-up of the participants.

**Procedures**

As a standard procedure, the participants received the same study questionnaires as used for the 1-year and 2-year follow-up every year. At approximately 5 years after the study inclusion, the participants were contacted once again by mail, but now with an accompanied letter and asked to fill out the study questionnaires as used for the 1 and 2 years’ follow-up with extra-questions about reoperations. Patients who did not respond initially were contacted by telephone by a research nurse and asked once again to participate in the study by filling out the questionnaires. The additional 5 years’ assessment was approved by the local medical ethics committee.

**Primary and secondary outcomes**

Primary outcome measures consisted of the Roland disability questionnaire (RDQ) for sciatica, a 100 mm visual analogue scale (VAS) for leg pain, and a seven-point Likert score of global perceived recovery. Higher RDQ and VAS scores were indicative of the experience of worse disability or greater intensity of pain, respectively. Global perceived recovery was measured with a seven-point Likert self-rating scale. Complete or almost complete disappearance of complaints (Likert scores 1–2) was defined as the ‘satisfactory recovery’, whereas Likert scores 3–7 were defined as ‘unsatisfactory recovery’.

Secondary outcomes were a 100 mm VAS for back pain and the number of (re)operations for severe sciatica in the interval between 2 and 5 years. The number of (re)operations for severe sciatica was evaluated by asking the patients whether there had been any new operations for severe sciatica in the interval between 2 and 5 years. Additionally, in all participating centres, it was checked whether patients had had a treatment because of sciatica in the intervening period.

**Potential prognostic factors**

The prognostic value of demographic and clinical baseline variables for unsatisfactory recovery at 5 years was evaluated. The initial list of prognostic factors, chosen in advance by the investigators, was based on potential clinical importance, as indicated by earlier clinical results. The following potential prognostic demographic variables were included in the analysis: age (dichotomised <40/≥40), gender, smoking status, body mass index (BMI) (dichotomised <25/≥25), physical job (yes/no) or mentally demanding job (yes/no).

The following clinical baseline variables were included in the analysis: level of herniation at the MRI, the presence of a sequester on MRI (yes/no), sciatica provoked by sitting (yes/no), sciatica provoked by coughing/sneezing (yes/no), outcome of Bragard’s test (positive/negative) and sensory disturbance (yes/no).

Furthermore, the following measurement instruments for general health, mental health, affective score and pain were included: the VAS scores for leg pain, back pain and general health, the Mental Health subscore of the Medical Outcomes Study 36-Item Short-Form General Health Survey (SF-36), and the McGill affective score. The VAS score range from 0 to 100, with...
higher scores indicating more severe symptoms. For this analysis, dichotomised VAS scores were used (<70/≥70), as described in an earlier study. The SF-36 Mental Health score ranges from 0 to 100, with higher scores indicating less severe symptoms. For this analysis, dichotomised SF-36 Mental Health subscores were used (scores below 1 SD of the Dutch reference population were defined as impaired). The McGill affective score measures the qualitative perception of pain by the patient and ranges from 0 to 5 where a high score (3–5) is correlated with a more depressed and anxious mood. For this analysis, dichotomised McGill affective scores were used (<3/≥3).

Statistical analysis
The original sample size calculation was based on a difference in RDQ outcome during and in a different speed to recovery during the first year. The main endpoint ‘recovery’ is in principle time-dependent in the sense that it reflects the situation of a patient at a particular moment in time and the situation may also deteriorate afterwards, so it can change from recovered to non-recovered and back to recovered again, therefore actually being a time-varying dichotomous outcome which should be taken into account when interpreting the results of the analyses. Differences between randomisation groups at baseline and after 5 years of follow-up were assessed by comparing means, medians or percentages, depending on the type of variable. Baseline values of variables were used as covariates in the main analyses whenever appropriate to adjust for possible differences between the randomised groups and to increase the power of the analyses. Outcomes of function and pain over the entire follow-up period were analysed using a repeated measurements analysis of variance with a first order autoregressive covariance matrix. Estimated consecutive scores were expressed as means and 95% CIs. Pointwise estimates were obtained using models with time

| Table 1 | Baseline and follow-up characteristics of patients with sciatica |
|---------|---------------------------------------------------------------|
| **Patient characteristics** | **Early surgery (n=141)** | **Conservative treatment (n=142)** |
| **Baseline characteristics** | | |
| Mean (SD) age (years) | 41.6 (10.0) | 43.3 (9.6) |
| Male sex | 89 (63) | 97 (68) |
| Mean (SD) BMI | 25.9 (4.1) | 25.5 (3.3) |
| Mean (SD) duration of sciatica (weeks) | 9.43 (2.37) | 9.48 (2.11) |
| Took sick leave from work | 107 (76) | 116 (82) |
| Mean (SD) duration of sick leave (weeks) | 5.32 (2.78) | 5.28 (2.62) |
| Left sided leg pain | 67 (48) | 73 (51) |
| Positive straight leg raising test (SLR) | 100 (71) | 104 (73) |
| Positive crossed SLR | 71 (50) | 70 (49) |
| Dermatomal sensory loss | 123 (87) | 128 (90) |
| Dermatomal anaesthesia | 31 (22) | 33 (23) |
| Dermatomal muscle weakness | 93 (66) | 99 (70) |
| Knee tendon reflex difference | 54 (38) | 51 (36) |
| Ankle tendon reflex difference | 75 (53) | 107 (75) |
| Clinically suspected level of herniated disc | | |
| L3–L4 | 6 (4) | 4 (3) |
| L4–L5 | 65 (46) | 52 (37) |
| L5–S1 | 70 (50) | 86 (61) |
| Preference for conservative treatment | 42 (30) | 43 (30) |
| Mean (SD) RD score | 16.5 (4.4) | 16.3 (3.9) |
| Mean (SD) VAS score | | |
| Leg pain | 67.3 (19.6) | 64.3 (21.2) |
| Back pain | 34.0 (29.6) | 30.8 (27.7) |
| Mean (SD) SF-36 scores | | |
| Bodily pain | 21.9 (16.6) | 23.9 (18.1) |
| Physical functioning | 33.9 (19.6) | 34.6 (19.0) |
| **Surgical treatment during follow-up** | | |
| Surgery performed in first year | 125 (89) | 55 (39) |
| Surgery performed during 2 years | 125 (89) | 62 (44) |
| Surgery performed during 5 years | 125 (89) | 66 (46) |
| Recurrent disc surgery after 5 years | 9 (6) | 8 (6) |
| Follow-up | 9 of 125 (7) | 8 of 66 (12) |
| Patients requiring two reoperations or more | 0 | 3 |
| Patients who dropped out during 5 year | 26 (18) | 26 (18) |

Values are numbers (percentages) unless stated otherwise.
BMI, body mass index; VAS, visual analogue scale.
as a categorical covariate to allow assessment of systematic patterns. Differences between groups in the dichotomised Likert score at 5 years were evaluated with Fisher’s exact test (randomisation group) or Mann-Whitney U (disability and pain scores).

The analyses were done according to the intention-to-treat principle, except the comparison of groups with a satisfactory or unsatisfactory recovery. Univariate and multivariate analyses were performed to evaluate the prognostic value of baseline variables for an unsatisfactory outcome at 5 years. The results are presented as ORs with 95% CIs. $\chi^2$ Tests were used to perform the univariate analyses. Potential predictors with a p<0.10 in the univariate analysis were included in the multivariate logistic regression. The multivariate logistic regression model was performed in a backward approach and included randomisation group irrespective of its significance in the univariate analysis, to control for its influence on the dependent variable.

For all other analyses, p<0.05 was considered significant. Data collection and quality checks were performed with the ProMISe web-based secure data management system of the Department of Medical Statistics and Bioinformatics of Leiden University Medical Centre. For all statistical analyses, SPSS V.18.0 was used.

RESULTS

Fifty-two of the 283 patients (18%) were lost to follow-up, among them one patient who died after a cardiac bypass operation and 19 patients who refused to participate. The baseline characteristics age, gender, BMI, randomisation group, RDQ and VAS-scores were not significantly different between the dropouts and those patients providing the 5 years’ follow-up data. Twenty-six of the dropouts were randomised for early surgery. Among the 26 dropouts in the prolonged conservative group, 11 (42%) had surgery for sciatica during follow-up. Among the 26 dropouts in the prolonged conservative group, 11 (42%) had surgery for sciatica during follow-up.

At 5 years’ follow-up, 66 of the 142 patients (46%) assigned to conservative treatment had had surgery because of intractable sciatica (table 1). Within the first year this was 55 (39%) and after 2 years 62 (44%). Of the 141 patients allocated for early surgery, 16 (11%) recovered before surgery and were not operated on during the 5 years of follow-up. Within this 5 years period, nine patients (7%) in the early surgery group and eight of the conservatively allocated patients who had surgery (12%) needed recurrent disc surgery. Three patients in the conservative group needed two reoperations or more (table 1).

The primary and secondary outcome scores concerning disability, leg pain and back pain at 5 years were slightly higher in the early surgery group compared with the prolonged conservative group; however, there were no significant differences (table 2 and figure 1A–C).

In the total group, after 5 years, 49 (21%) patients still had an unsatisfactory recovery, defined as not having a

| Outcomes | 8 weeks | 52 weeks | 130 weeks |
|----------|---------|----------|-----------|
| Disability* | ES | PCT | Difference (95% CI) |
| Leg pain† | ES | PCT | Difference (95% CI) |
| Back | ES | PCT | Difference (95% CI) |

*Roland disability questionnaire for sciatica. Score ranges from 0 to 23, with higher scores representing worse disability.
†Measured on a 100 mm visual analogue scale, with 0 representing no pain and 100 the worst pain ever experienced.
Values are means (SE) unless stated otherwise and are based on intention to treat, CI assessed with repeated measurements analysis.
complete or almost complete recovery on the dichotomised Likert scale, irrespective of their allocated treatment group (25 patients in the prolonged conservative group, 24 patients in the early surgery group, p=1.00).

Patients with an unsatisfactory recovery had a significantly higher amount of leg pain, back pain and disability (all p values <0.01) compared with the group with a satisfactory recovery (table 3). At 5 years' follow-up, 93 patients (62 men and 31 women, mean age 43.7, SD 9.8) had not had surgery (76 from the conservative group and 16 from the early surgery group) and the percentage recovered (77%) or not-recovered at 5 years (23%) was not different from the total group.

In the period of 5 years, 66 of the 213 patients (31%) with collected data at one, 2 and 5 years of follow-up had at least one period of unsatisfactory recovery. The pattern of recovery showed 16 patients who did not report any recovery at 1, 2 and 5 years of follow-up. Twenty-four patients switched from ‘not recovered’ in the first or second year to ‘recovered’ at the 5 years’ analysis. Sixteen of these patients were from the prolonged conservative group. Six of these patients needed an operation of whom four needed a reoperation before they recovered, compared with two of the eight patients in the early surgery group needing a reoperation. Twenty-six patients showed a good recovery at 1 and/or 2 years but not at 5 years of follow-up (figure 2).

Univariate logistic regression evaluating the relationship between possible prognostic variables and unsatisfactory recovery at 5 years, irrespective of intermediate recovery, showed that a high McGill affective score (score >3; OR 6.23 (95% CI 2.23 to 17.38)) was a significant predictor, as were severe leg pain at baseline (VAS >70; OR 2.95 (95% CI 1.54 to 5.64)), an impaired score on the SF-36 Mental Health subscale (OR 2.25 (95% CI 1.16 to 4.35)), higher age (age >40; OR 2.22 (95% CI 1.11 to 4.47)) and a positive Bragard test (OR 1.91 (95% CI 0.97 to 3.74); table 4). The multivariate analysis included all variables that were related to unsatisfactory outcome in the univariate analysis (p<0.10), as well as randomisation group. This analysis showed that a high McGill affective score (score >3; OR 4.48 (95% CI 1.43 to 14.08)), severity of leg pain (VAS >70; OR 2.80 (95% CI 1.39 to 5.62)) and age (age >40 years; OR 2.36 (95% CI 1.12 to 5.00)) were the only significant predictors for an unsatisfactory outcome at 5 years.

**DISCUSSION**

This long-term follow-up study of the same patient cohort corroborates with earlier 1-year and 2-year results as no significant differences between randomisation groups in disability, leg pain or back pain are found after 5 years of follow-up.2,3 Eighteen per cent of the initial cohort of 283 patients was lost to follow-up after 5 years. This reduced the power of our latest analysis to some extent. However, the baseline characteristics showed no differences between the included group and the dropouts.

The allocated strategy of an extra 6 months of wait-and-see resulted in a large proportion of delayed surgical treatment (46%) for persistent intense leg pain.
causing severe disability, despite all kinds of conservative treatment. This means that patients should be informed that prolonged conservative care gives them quite a high chance for resolution of pain and disability without the need of a surgical intervention, but that this strategy also carries a fair chance (46%) that this waiting for the pain to resolve will still end with them needing disc surgery.

The question, however, is whether this conclusion is completely accurate. The design of the original study was a comparison of early surgery versus prolonged conservative care for 6 months, after which the surgery was offered when there still were severe complaints. This means that we do not know the precise percentage of patients who would have become pain free with even longer conservative care, because the majority of these patients were operated on after 6 months. But, on the other hand, one might question whether such a proposed long-conservative regimen is in proportion with the small risk of a surgical intervention, which provides a better outcome in the first 6 months, rather than being disabled in daily life during this long period of conservative care. Furthermore, the proportion of patients who were not operated after 5 years with a satisfactory or unsatisfactory recovery was the same as in the total population, showing that also in the non-operated patients there was a rather high amount of unsatisfactory recovery.

Twenty-one per cent of patients experienced unsatisfactory recovery at 5 years, while 31% of the patients with complete data at 1, 2 and 5 years of follow-up noted at least once an unsatisfactory recovery during this 5 years’ follow-up period, irrespective of their allocated group. So, the optimal timing of surgery is still on debate, as is the question which patients would benefit from surgery and which from prolonged conservative care.

The study design of this 5 year analysis did not permit us to look properly for the causes of an unsatisfactory recovery. A shortcoming of this study is the fact that there was no permission in the present study to retrieve new MRI, but a recent study with the same patient population did not show any correlation between MRI and satisfactory or unsatisfactory recovery at 1 year. The patterns of recovery show that 8% of all patients with collected data had never had any recovery during the follow-up period, showing that there are not that many non-responders of conservative or surgical treatment. The other 50 patients (25%) with an unsatisfactory recovery showed a switch over time from recovered to not-recovered, or vice versa. This is in line with the idea that sciatica is caused by chronic disc disease with intermittent nerve compression or inflammation and that pain can thus reoccur despite any earlier treatment. In the 24 patients who switched from ‘not-recovered’ in the first or second year to ‘recovered’ at the 5-years’ analysis, there was a higher amount of patients who needed a reoperation before recovery occurred in the prolonged conservative group compared with the early surgery group. This may be caused by less effectiveness of late surgery, although this could not be

| Outcome                   | Disability* Mean (SD) | Median (IQR) | Leg pain† Mean (SD) | Median (IQR) | Back pain† Mean (SD) | Median (IQR) |
|---------------------------|-----------------------|--------------|---------------------|--------------|---------------------|--------------|
| Unsatisfactory recovery   | 11.3 (5.6)            | 1.0 (6.5–15.5)| 42.8 (27.9)         | 46.0 (17.5–66.0)| 48.6 (49.5)         | 49.5 (32.0–66.75) |
| (n=49)‡                   | 1.3 (2.3)             | 0.0 (0.0–2.0) | 6.5 (13.7)          | 1.0 (0.0–6.0)  | 10.2 (14.6)         | 4.0 (0.0–15.0)  |

*Roland disability questionnaire for sciatica. †Measured on a 100 mm analogue scale. ‡The seven-point Likert scale of global perceived recovery was dichotomised to satisfactory outcome (‘complete’ and ‘nearly complete’ recovery) and unsatisfactory outcome (the other 5 scores ranging from ‘some recovery’ to ‘severe worsening of complaints’).
| Variable                                      | Univariate analysis | Multivariate analysis |
|-----------------------------------------------|---------------------|-----------------------|
|                                               | N       | OR  | 95% CI     | p Value | OR  | 95% CI     | p Value |
| Randomisation                                 |         |     |            |         |     |            |         |
| Surgery                                       | 115     | 0.96| 0.51 to 1.80 | 0.899   |     |            |         |
| Conservative                                  | 116     | 1.00| –           |         |     |            |         |
| Gender                                        |         |     |            |         |     |            |         |
| Female                                        | 75      | 1.27| 0.66 to 2.46 | 0.472   |     |            |         |
| Male                                          | 156     | 1.00| –           |         |     |            |         |
| Age                                           |         |     |            |         |     |            |         |
| ≥40                                           | 137     | 2.22| 1.11 to 4.47 | 0.023   | 2.36| 1.11 to 5.00 | 0.024   |
| <40                                           | 94      | 1.00| –           |         |     |            |         |
| Mentally demanding job                        |         |     |            |         |     |            |         |
| Yes                                           | 139     | 1.17| 0.57 to 2.37 | 0.669   |     |            |         |
| No                                            | 76      | 1.00| –           |         |     |            |         |
| Physical job                                  |         |     |            |         |     |            |         |
| Yes                                           | 85      | 1.39| 0.71 to 2.70 | 0.339   |     |            |         |
| No                                            | 132     | 1.00| –           |         |     |            |         |
| Smoking                                       |         |     |            |         |     |            |         |
| Yes                                           | 84      | 1.09| 0.57 to 2.09 | 0.792   |     |            |         |
| No                                            | 142     | 1.00| –           |         |     |            |         |
| BMI                                           |         |     |            |         |     |            |         |
| ≥25                                           | 125     | 1.19| 0.62 to 2.29 | 0.605   |     |            |         |
| <25                                           | 101     | 1.00| –           |         |     |            |         |
| Sciatica provoked by sitting                  |         |     |            |         |     |            |         |
| Yes                                           | 180     | 0.73| 0.35 to 1.51 | 0.397   |     |            |         |
| No                                            | 51      | 1.00| –           |         |     |            |         |
| Coughing, sneezing                           |         |     |            |         |     |            |         |
| Yes                                           | 163     | 1.06| 0.53 to 2.12 | 0.881   |     |            |         |
| No                                            | 68      | 1.00| –           |         |     |            |         |
| Bragard’s test                                |         |     |            |         |     |            |         |
| Positive                                      | 62      | 1.91| 0.97 to 3.74 | 0.059   |     |            |         |
| Negative                                      | 154     | 1.00| –           |         |     |            |         |
| Sensory disturbance                           |         |     |            |         |     |            |         |
| Yes                                           | 200     | 1.45| 0.47 to 4.47 | 0.514   |     |            |         |
| No                                            | 24      | 1.00| –           |         |     |            |         |
| VAS leg pain                                  |         |     |            |         |     |            |         |
| ≥70                                           | 89      | 2.95| 1.54 to 5.64 | 0.001   | 2.80| 1.39 to 5.62 | 0.004   |
| <70                                           | 142     | 1.00| –           |         |     |            |         |
| VAS back pain                                 |         |     |            |         |     |            |         |
| ≥70                                           | 34      | 1.68| 0.74 to 3.80 | 0.211   |     |            |         |
| <70                                           | 196     | 1.00| –           |         |     |            |         |
| McGill affective score                        |         |     |            |         |     |            |         |
| High (3–5)                                    | 17      | 6.227| 2.23 to 17.38 | <0.001  | 4.48| 1.43 to 14.08 | 0.010   |
| Low (0–2)                                     | 209     | 1.00| –           |         |     |            |         |
| MRI-level herniation                          |         |     |            |         |     |            |         |
| L5S1                                          | 125     | 0.69| 0.37 to 1.31 | 0.256   |     |            |         |
| L3L4/L4/L5                                    | 106     | 1.00| –           |         |     |            |         |
| MRI-sequester                                 |         |     |            |         |     |            |         |
| Yes                                           | 84      | 0.85| 0.43 to 1.68 | 0.643   |     |            |         |
| No                                            | 122     | 1.00| –           |         |     |            |         |
| SF-36 mental health*                          |         |     |            |         |     |            |         |
| Impaired                                      | 66      | 2.25| 1.16 to 4.35 | 0.014   |     |            |         |
| Not impaired                                  | 163     | 1.00| –           |         |     |            |         |
| VAS general health                            |         |     |            |         |     |            |         |
| ≥70                                           | 60      | 1.36| 0.68 to 2.74 | 0.382   |     |            |         |
| <70                                           | 168     | 1.00| –           |         |     |            |         |

Only the numbers of the potential predictors with a p < 0.10 in the univariate analysis were shown in the multivariate logistic regression.

*The scores on the SF-36 Mental Health subscale were dichotomised using 1 SD below the Dutch reference population.

BMI, body mass index; VAS, visual analogue scale.
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proven. This less effectiveness of late surgery compared with early surgery could be caused by more chronic changes around the disc protrusion or sequester, causing more difficulty in freeing the nerve from compression.

In our study, we could not estimate the effect of early versus late surgery, or surgery versus conservative treatment in an unbiased way since that would require a per protocol analysis. This would ignore the randomised allocation and compare patients who by definition would be incomparable since the design of our study did not envisage a randomisation between early and late surgery, but merely allowed for the time of surgery being determined by the assessment of both patient and physician after initial randomisation. The impossibility to capture completely the condition of the patient and the selection mechanism which leads to the decision to operate, or not, at any given point in (follow-up) time, renders any multivariate analysis that attempts to make the groups of early and late operations comparable, biased. This bias has occurred in other randomised studies where this ‘per protocol’ analysis was used and patients who were operated on after a prolonged duration of symptoms from a herniated lumbar disk have been shown to have a worse outcome than patients who were operated on relatively early. As a consequence of this E “per protocol analysis both groups differed in baseline characteristics such as type of disc herniation, neurological deficit and reported depression, rendering comparison of both groups fallacious.”

In this 5-year analysis the independent prognostic factors for an unsatisfactory outcome were a high McGill affective score, a high amount of leg pain and age over 40 years at baseline. A high McGill affective score correlates with a more depressed and anxious mood, and mental stress, depression or other psychological factors have been widely described as risk factors for development of chronic pain. Although a high amount of pain at baseline can also be caused by psychological factors, it is an independent prognostic factor for an unsatisfactory outcome in this study. Perhaps the severity of nerve root compression at intake predicts the overall outcome, but a correlation between amount of pain and amount of root compression has not been proven. In our study, other factors concerning the severity of nerve root damage, such as severe sensibility disturbance, showed no association. In a systematic review of non-surgically treated sciatica, none of the three studies that investigated baseline leg pain severity showed a clear prognostic influence on outcome. Age was also not found to be a prognostic factor in six of seven studies. The reason for age over 40 years being a prognostic indicator for an unsatisfactory outcome could be because disc herniation is part of a degenerative disease which seems to worsen in time. A study of prognostic factors for unsatisfactory recovery among operated and unoperated patients did not show leg pain severity and age to be a prognostic factor, but found the variables severe back pain and male gender as predictors for a bad outcome at the 1-year follow-up. This was in contrast with our 1-year study where being woman was a prognostic factor for a bad outcome, but in this 5 years’ analysis this also disappeared. The possible prognostic role of gender might be overestimated in the past.

CONCLUSION

After 5 years of follow-up, there were still no differences in pain and disability between the patients randomised for early surgery or prolonged conservative care.

Signs of pain quality associated with depression and a more anxious mood, the age of the patient and the severity of leg pain at baseline were predictive of an unsatisfactory outcome.

In general, patients must be informed that prolonged conservative care might give them a fair chance for pain and disability to resolve without surgery, but with the risk for delayed surgery in the end after a prolonged period of suffering from sciatica.

Furthermore, although the total number of disabled patients without pain free periods in this study seems to be low, in almost one-fourth of the patients, sciatica appears to be an ongoing disease with variable complaints in time, irrespective of treatment. These patients need our full attention, and further investigation at 10 years’ follow-up is being planned, because this patient category has the highest burden on society concerning work absenteeism and general health costs.

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Contributors WCP, RB, GJB and MBL conceived the idea of this follow-up study and were responsible for the design of the study. MBL, DV, WCHJ and RB were responsible for collecting the data, data analysis and MBI, DV and RB were responsible for tables and graphs. WCP, WPV, GJB and WCHJ provided input in the data analysis. The initial draft of the manuscript was prepared by MBL and DV and then circulated repeatedly among all authors for
critical revision. MBL, DV, WCHJ and RB were responsible for the acquisition of the data and WCP, WPV, GJB, RB, DV and MBL contributed to the interpretation of the results. All authors have read and approved the final manuscript.

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