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BACKGROUND
Leprosy causes nerve damage which may result in nerve function impairment and disability. Decompressive surgery is used for treating nerve damage, although the effect is uncertain.

OBJECTIVES
To assess the effects of decompressive surgery on nerve damage in leprosy.

SEARCH STRATEGY
We searched the Cochrane Neuromuscular Disease Group Trials Register (November 2007), the Cochrane Central Register of Controlled Trials (The Cochrane Library Issue 4, 2007), MEDLINE (from January 1950 to November 2007), EMBASE (from January 1980 to November 2007), AMED (from January 1985 to November 2007), CINAHL (from January 1982 to November 2007) and LILACS (from January 1982 to November 2007) in November 2007. We checked reference lists of the studies identified, the Current Controlled Trials Register (www.controlled-trials.com), conference proceedings and contacted trial authors.

SELECTION CRITERIA
Randomised and quasi-randomised controlled trials of decompressive surgery for nerve damage in leprosy.

DATA COLLECTION AND ANALYSIS

The primary outcome was improvement in sensory and motor nerve function after one year. Secondary outcomes were improvement in nerve function after two years, change in nerve pain and tenderness, and adverse events. Two authors independently extracted data and assessed trial quality. We contacted trial authors for additional information. We collected adverse effects information from the trials and non-randomised studies.

MAIN RESULTS
We included two randomised controlled trials involving 88 people. The trials examined the added benefit of surgery over prednisolone for treatment of nerve damage of less than six months duration. After two years follow-up there was no significant difference in improvement in nerve function between surgery and prednisolone treatment.
nerve function improvement between people treated with surgery plus prednisolone or with prednisolone alone. Adverse effects of decompressive surgery were not adequately described.

**Authors' conclusions**

Decompressive surgery is used for treating nerve damage in leprosy but evidence from randomised controlled trials does not show a significant added benefit of surgery over steroid treatment alone. Well-designed randomised controlled trials are needed to establish the effectiveness of the combination of surgery and medical treatment compared to medical treatment alone.

**PLAIN LANGUAGE SUMMARY**

**Decompressive surgery for treating nerve damage in leprosy**

Leprosy is a chronic infectious disease. Leprosy bacteria cause damage to skin and peripheral nerves which may result in nerve function impairment and disability. Decompressive surgery is used for treating nerve damage although its effect is uncertain. Two randomised controlled trials were included in the review and examined the added benefit of surgery over prednisolone for treatment of nerve damage of less than six months duration. Two years from the start there was no significant difference in nerve function improvement between people treated with surgery plus prednisolone or with prednisolone alone. Adverse effects of decompressive surgery were not adequately described.
**Description of the condition**

Leprosy is a chronic infectious disease caused by the bacillus *Mycobacterium leprae*. Leprosy bacilli are spread in tiny droplets from the nose or mouth from infected and untreated individuals. When the immune system fails to respond effectively to the antigens of the bacilli, the disease will develop. Leprosy bacilli may directly or indirectly cause damage. Often, the first sign of leprosy is a skin lesion. Damage to peripheral nerves may cause symptoms such as loss of sweating, sensation and muscle strength. Leprosy appears in various clinical forms, dependent on the response of the immune system. Some people have only a few skin lesions and the number of bacilli is relatively small. This is classified as paucibacillary (PB) leprosy. Other people may have many skin lesions with multiple nerves involved and a high number of bacilli in their body and are then classified as having multibacillary (MB) leprosy (ILEP 2001; WHO 2006).

Leprosy can be effectively treated with a combination of antibiotics (rifampicin, dapsone and clofazimine). Since the introduction of this multidrug therapy (MDT), the number of people with leprosy has decreased substantially. At the beginning of 2007 the prevalence was about 225,000 worldwide. This is the registered number of people on MDT treatment. The number of newly detected people reported was approximately 259,000 in 2006 (WHO 2007).

**Causes**

The body’s immune response to the antigens of the leprosy bacilli may cause periods of inflammation in the skin and peripheral nerves and sometimes also in other organs: so-called ‘reactions’. There are two types of potentially nerve damaging reactions: type 1 reaction or reversal reaction (RR) and type 2 reaction or erythema nodosum leprosum (ENL). Reactions may occur before, during and after multidrug therapy and are the main cause of nerve damage and consequently of impairment in leprosy (ILEP 2002; Lockwood 2005; WHO 1998). Nerve damage may develop slowly and is often unnoticed until very late. It is often the symptoms of a reaction that force people to seek medical help (Job 1989; Nicholls 2003).

**Impact**

Leprosy is a most important disabling disease. The World Health Organization estimated the number of people living with physical disabilities due to leprosy at two to three million worldwide (WHO 2004). Despite a fast declining trend in the number of newly detected people with leprosy, the decline in the number of people living with physical disabilities is much slower. In the near future, we can still expect about one million people to be affected by leprosy disability (Meima 2004). People affected by leprosy, especially those with visible deformities and disabilities, fear discrimination and stigmatisation. These people may experience severe social and psychological problems (Heijnders 2004; Leekassa 2004; Rafferty 2005).

**Treatment**

Corticosteroids, especially prednisolone, are used as first-line treatment of severe reversal reactions and nerve damage in leprosy. They work by controlling the acute inflammation and relieving the pain (Britton 1998; Lockwood 2000). The earlier corticosteroids are given after the onset of nerve damage, the more likely permanent nerve function impairment will be prevented (Becx-Bleumink 1990; Naafs 1996). Prednisolone seems to be a very effective drug, but it has some shortcomings. Long-term therapy may cause serious adverse effects, such as peptic ulcer, cataract or psychosis (Richardus 2003; Sugumaran 1998; WHO 1998). A considerable proportion of people treated for severe reversal reactions and nerve damage does not benefit from standard corticosteroid treatment (Croft 2000; Lockwood 1993; Saunderson 2000; Schreuder 1998). The long-term benefit of corticosteroids for treating nerve damage is still uncertain and trials establishing the optimal regimen and effectiveness are needed (Van Veen 2007).

Other immunosuppressant drugs for treating severe reversal reactions and nerve damage have been tested or are under examination, such as azathioprine (Marlowe 2004) and cyclosporin A (Lockwood 2000; Sena 2006). It is plausible that these drugs may be effective for treating nerve damage, but evidence from randomised controlled trials is very limited.

Decompressive surgery or neurolysis as treatment for nerve damage has been used for several decades. The objective of this surgery is to relieve mechanical compression, due to oedema caused by neuritis, of the affected nerve. Decompression is done by incision of the thickened nerve sheath (epineurium) where the nerve is enlarged and often tender on palpation. This incision is often of a considerable length at the place before entering the fibrous tunnel which, during surgery, also needs to be opened. Results from non-randomised studies of surgery have been widely published (e.g. Brandsma 1983; Carayon 1993; Chaise 1985; Dandapat 1991; Droogenbroeck 1977; Ramarozaraza 1995; Rao 1989), but there is little evidence from randomised controlled trials that surgery is better than medical treatment alone (Boucher 1999; Ebenezer 1996; Pannikar 1984). Decompressive surgery is not recommended without medical treatment. Indications for surgery are mainly based on common practice but not well-defined. These may include the presence of nerve abscess, nerve pain or nerve function impairment that does not respond to medical treatment (Chaise 2004; Kazen 1996; Malaviya 2004b; Richard 2004; Palande 1980).

**Why it is important to do this review**

Decompressive surgery is frequently used for treating nerve damage in leprosy. The effect of surgery, especially in the long-term, is uncertain and it is unclear whether surgery is more beneficial...
than medical treatment alone. While this review focused on evidence from randomised controlled trials (RCTs), it was expected that only a few RCTs would have been conducted in this area. Therefore, the results were also considered in the light of non-randomised evidence in the Discussion section.

**OBJECTIVES**

To assess the effects of decompressive surgery for treating nerve damage in leprosy.

**METHODS**

**Criteria for considering studies for this review**

**Types of studies**

All randomised controlled trials (RCTs) and quasi-randomised controlled trials (quasi-RCTs) of any design.

**Types of participants**

Anyone with leprosy confirmed by appropriate clinical signs or symptoms according to Ridley 1966 or WHO 1998 classification and leprosy-related nerve damage or severe leprosy type 1 reaction, requiring corticosteroid treatment. Nerve damage or nerve function impairment was defined as clinically detectable impairment of motor or sensory nerve function. It did not include impairment of nerve conduction that was only detectable by electrophysiological means (Croft 1999).

**Types of interventions**

Decompressive surgery or neurolysis for treating nerve damage in leprosy. The comparators were no treatment, placebo or corticosteroids.

**Types of outcome measures**

**Primary outcomes**

(1) (i) Improvement in sensory nerve function one year after registration, as determined and defined by the original authors. Sensory nerve function was assessed with five graded nylon filaments or a ball-point pen. We adapted the scores as defined by Van Brakel et al (Van Brakel 2005). For testing with graded nylon filaments, sensory nerve function impairment was diagnosed if the monofilament threshold was increased from normal by three or more points for any nerve. One point was given for each level that the monofilament threshold was increased from normal at each test site. The points were added for each nerve. Normal thresholds used were 200 milligrams for the hand and two grams for the foot. If the score for any nerve decreased by three or more points from the baseline score, the nerve was considered as improved. When a non-graded test was used, such as the ball-point pen test, a nerve was diagnosed as impaired if two or more test sites did not feel the stimulus. Improvement for any nerve was defined as two or more test sites feeling the stimulus, compared to the baseline measurement (Van Brakel 2003). Improvement was a dichotomous outcome variable (improvement or not).

(ii) Improvement in motor nerve function one year after registration, as determined and defined by the original authors. Improvement in motor nerve function was assessed with the modified MRC grading scale (Brandsma 1981). Motor nerve function impairment was defined as a score of less than four on the modified MRC grading scale for any nerve. Improvement was defined as at least one point improvement in score for any muscle compared to the initial score. Improvement was a dichotomous outcome variable (improvement or not).

(2) Adverse outcomes

We documented the incidence and severity of all recorded local and systemic adverse events, at any time point, in all the included studies.

(3) Economic data

We did not report data relating to costs but we addressed cost implications in the Discussion section if information was available.

(4) Timing of outcome assessment

Data that have been recorded for less than six months were considered to reflect short-term benefit and were analysed separately from data that were recorded after one year or more, which we considered to reflect the minimum time period to capture any long-term benefit. The end point closest to three months (one to six months) was used for short-term benefit and the end point closest to two years (one year) was used for long-term benefit. The long-term data were considered the primary endpoint but we considered the short-term data in order to detect rapid onset of improvement.

**Search methods for identification of studies**

**Electronic searches**

We searched for relevant published trials in the Cochrane Neuromuscular Disease Group Trials Register (November 2007) using the following terms: leprosy or Hansen disease and decompression or neurolysis or epicondylectomy or epineurotomy or neuritis or nerve damage or nerve loss or nerve function impairment or neuropathy or nerve problem or nerve involvement or nerve
pain. This search strategy was adapted to include additional search terms where necessary and was modified to search the Cochrane Central Register of Controlled Trials in *The Cochrane Library* (Issue 4, 2007); MEDLINE (from January 1950 to November 2007) and EMBASE (from January 1980 to November 2007); AMED (Allied and Complementary Medicine, from January 1985 to November 2007), CINAHL (from January 1982 to November 2007), and LILACS (Latin American and Caribbean Health Science Information database, from January 1982 to November 2007). See Appendix 1, Appendix 2, Appendix 3, Appendix 4 and Appendix 5 for MEDLINE, EMBASE, AMED, LILACS and CINAHL strategies respectively. We searched for ongoing trials in the metaRegister of Controlled Trials (www.controlled-trials.com).

**Searching other resources**

**Reference lists**

We scanned the bibliographies of the included studies and reviews for possible references to RCTs.

**Unpublished literature**

We attempted to find unpublished or ongoing trials via correspondence with trial authors of included and excluded trials less than 15 years old and other disease experts.

**Handsearching**

Conference proceedings from relevant leprosy meetings were scanned for RCTs and, where possible, the authors were contacted for further information.

**Adverse effects**

We did not perform a separate search for adverse events.

**Language restrictions**

No language restrictions were imposed when searching for publications, and translations were sought where necessary.

**Data collection and analysis**

**Selection of studies**

Two review authors (NvV and JHR) checked the titles and abstracts identified from the searches. If it was clear that the study did not refer to a randomised controlled trial of surgical decompression for treating nerve damage in leprosy, we excluded it. The same two review authors independently assessed the full text version of each remaining study to determine whether it met the predefined selection criteria. Any differences of opinion were resolved through discussion within the review team. We listed the excluded studies and reasons for exclusion in the 'Characteristics of excluded studies' table.

**Data extraction and management**

Two review authors (NvV and JHR) independently extracted data from the included studies onto a data extraction form. If there were missing data, we contacted the trial authors. We entered data into Review Manager (RevMan) (RevMan 2008). Authors were not blinded to trial author, journal or institution.

We were not able to translate reported changes in nerve function and nerve pain into the proportion of participants with improvement greater than minimal. By minimal we meant anything greater than 50% improvement from baseline on a continuous scale for sensory nerve function or a score of four or more on the MRC grading scale for motor nerve function. It was not possible to calculate the proportion of participants with full recovery of nerve function.

**Assessment of methodological quality**

The quality assessment included an evaluation of the following components for each included study, since there is some evidence that these are associated with biased estimates of treatment effect (Juni 2001):

(a) the method of generation of the randomisation sequence;
(b) the method of allocation concealment - it was considered 'adequate' if the assignment could not be foreseen;
(c) whether the outcome assessor was blinded;
(d) how many participants were lost to follow up in each arm, and whether reasons for losses were adequately reported;
(e) whether all participants were analysed in the groups to which they were originally randomised (intention-to-treat principle).

In addition we reported on:

(f) the baseline assessment of the participants for age, sex, duration and severity of nerve function impairment;

(g) whether outcome measures were described.

We described the quality of each study, based on these components, in the section on Risk of bias in included studies. We also recorded this information in Table 1.
Table 1. Methodological quality assessment

<table>
<thead>
<tr>
<th>Study ID</th>
<th>Allocation concealment</th>
<th>Patient blinding</th>
<th>Assessor blinding</th>
<th>Loss to follow-up</th>
<th>Clear diagnosis</th>
<th>Baseline differences</th>
<th>Explicit outcomes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Boucher 1999</td>
<td>A: computer random number table</td>
<td>C: not possible</td>
<td>C: not possible</td>
<td>B: unclear for nerves A: 2/31 participants (6%)</td>
<td>A</td>
<td>A</td>
<td>A</td>
</tr>
<tr>
<td>Pannikar 1984</td>
<td>C: alternation</td>
<td>C: not possible</td>
<td>C: not possible</td>
<td>C: 13/75 nerves (17%), 13/57 participants (23%)</td>
<td>A</td>
<td>A</td>
<td>A</td>
</tr>
<tr>
<td>Ebenezer 1996</td>
<td>C: alternation</td>
<td>C: not possible</td>
<td>C: not possible</td>
<td>C: 18/75 nerves (24%), 18/57 participants (32%)</td>
<td>A</td>
<td>A</td>
<td>A</td>
</tr>
</tbody>
</table>

**Measures of treatment effect**

We used the Cochrane statistical package, RevMan (RevMan 2008) for statistical data analysis. None of the study results could be pooled meaning that no weighted treatment effect could be calculated. Results were expressed as mean differences with 95% confidence intervals (CI) for continuous outcome measures and relative risks (RR) with 95% CI for dichotomous outcomes. One trial reported median improvement as outcome. It was not possible to perform tests for heterogeneity or sensitivity analysis due to insufficient trials. We will perform such analyses should trials become available in the future.

**Dealing with missing data**

We were not able to conduct an intention-to-treat analysis. By contacting the authors we learned that information about the groups to which lost participants were randomised was no longer available.

**Adverse outcomes**

In our Discussion, we considered adverse effects by taking non-randomised literature into account, since randomised studies rarely capture adverse events adequately.

**Economic issues**

We were not able to consider the costs and cost-effectiveness of treatment due to lack of evidence.

**RESULTS**

**Description of studies**

See: Characteristics of included studies; Characteristics of excluded studies.

We identified 10 potentially relevant studies and excluded seven, because they were not randomised. For a description of excluded trials see the ‘Characteristics of excluded studies’ table.

Two RCTs (one RCT was described in two papers), involving 88 people, were included. Both tested decompression surgery plus oral corticosteroids versus oral corticosteroids alone. One tested treatment of ulnar neuritis of less than six months duration (Ebenezer 1996; Pannikar 1984) and one tested treatment of neuritis of several types of less than six months duration (Boucher 1999).

For a full description of included trials, see the ‘Characteristics of included studies’ table.
The methodological quality assessment took into account concealment of allocation, patient blinding, outcome assessor blinding, completeness of follow-up, explicit diagnostic criteria, how studies dealt with baseline differences of the experimental groups, and explicit outcome criteria. We each graded these items: A: adequate, B: unclear, C: inadequate. If the information was not available the item was graded ‘inadequate’. The scores of each trial for all the quality measures are described in Table 1.

**Allocation**
In one trial (Boucher 1999) participants were randomly assigned to either intervention or control group and the allocation concealment was considered adequate. The other trial (Ebenezer 1996; Pannikar 1984) used alternation as the randomisation procedure which was considered inadequate.

**Blinding**
Participant and outcome assessor blinding was not possible in any of the trials.

**Follow-up**
One trial (Boucher 1999) had 6% loss to follow-up of participants, but did not report how many nerves were involved. The other trial (Ebenezer 1996; Pannikar 1984) had 17% loss to follow-up of nerves after one year and 24% loss to follow-up of nerves after two years. None of the trials reported how many participants or nerves were lost to follow up in each arm. Boucher et al described the reasons for losses.

**Diagnostic criteria**
Both trials diagnosed and classified leprosy using the internationally accepted diagnostic criteria of Ridley and Jopling (Ridley 1966).

**Baseline differences**
In the trials the baseline characteristics in both arms were similar.

**Explicit outcomes**
The primary outcomes 'improvement in sensory nerve function one year after registration' and 'improvement in motor nerve function one year after registration' were evaluated in one trial (Pannikar 1984). The secondary outcome 'improvement in nerve function two years after registration' was evaluated in two trials (Boucher 1999; Ebenezer 1996). 'Change in nerve pain and in nerve tenderness' was assessed in one trial (Pannikar 1984) one year after registration and in two trials (Boucher 1999; Ebenezer 1996) two years after registration. None of the trials evaluated 'changes in quality of life'. Adverse events were not well-reported in any of the trials.

**Effects of interventions**

**Medial epicondylectomy and nerve decompression plus oral corticosteroids versus oral corticosteroids alone for participants with ulnar neuritis of less than six months duration (Pannikar 1984; Ebenezer 1996)**

**Primary outcome measures**

(1) Improvement in sensory nerve function one year after registration
The trial compared oral prednisolone plus medial epicondylectomy and external nerve decompression (surgery group) with oral prednisolone alone (medical group) in participants with ulnar neuritis of less than six months duration (n = 57 participants with 75 nerves). One year after admission results of sensory nerve function were available for 31 nerves in the surgery group and 31 nerves in the medical group. Improvement was measured as either a mean change score between baseline and end of follow-up or the proportion of nerves with improvement. Sensory testing was done with a No.3 and No.6 nylon thread (approximately 200 mg and 5 g, respectively). Fifteen sites on the ulnar nerve distribution area were tested. The score for each nerve depended on the number of sites felt. The score was 15 when all sites were felt with the 200 mg thread, and zero when no site was felt with either thread. Sensory improvement was defined as a positive difference between the final and initial score. Mean differences between the baseline score and the score at the end of one year were compared for the two treatment groups. Results were available for 29 nerves in the surgery group and 28 nerves in the medical group. After one year the mean difference was 2.08 (95% CI 0.28 to 3.88) in the surgery group and 2.00 (95% CI 0.06 to 3.94) in the medical group, indicating a mean improvement in both. The improvement was slightly greater in the surgery group but the mean difference 0.08 (95% CI -2.45 to 2.61) between the two groups was not significant (Figure 1; Analysis 1.1). In the surgery group 18 out of 31 nerves (58%) had sensory improvement compared with 16 out of 31 nerves (52%) in the medical group. The difference was not significant (relative risk 1.13, 95% CI 0.71 to 1.77) (Figure 2; Analysis 1.2).
Figure 1. Forest plot of comparison: 1 Medial epicondylectomy plus oral steroids versus oral steroids alone, outcome: 1.1 Change in sensory score after one year.

<table>
<thead>
<tr>
<th>Study or Subgroup</th>
<th>Treatment Mean</th>
<th>SD</th>
<th>Total</th>
<th>Control Mean</th>
<th>SD</th>
<th>Total</th>
<th>Weight</th>
<th>Mean Difference</th>
<th>IV, Fixed, 95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Eterneeder 1996</td>
<td>2.08</td>
<td>4.74</td>
<td>28</td>
<td>2</td>
<td>5</td>
<td>28</td>
<td>100.0%</td>
<td>0.08 [-2.45, 2.61]</td>
<td></td>
</tr>
<tr>
<td>Total (95% CI)</td>
<td>29</td>
<td>28</td>
<td>100.0%</td>
<td>0.08 [-2.45, 2.61]</td>
<td></td>
<td></td>
<td></td>
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</tr>
</tbody>
</table>

Heterogeneity: Not applicable
Test for overall effect: Z = 0.86 (P = 0.39)
(2) Improvement in motor nerve function one year after registration

The trial measured motor improvement of the ulnar nerve at one year from the time of admission (n = 57 participants with 75 nerves). Results of motor nerve function were available for 31 nerves in the surgery group and 31 nerves in the medical group. Improvement was measured as either a change score between baseline and end of follow up or as the proportion of nerves improved. Motor nerve function of the ulnar nerve was assessed with the MRC grading scale. The maximum score for each muscle was five and for the whole nerve 15. Motor improvement was defined as a positive difference between the final and initial score. Mean differences between the baseline score and the score at the end of one year were compared for the two treatment groups. Results were available for 29 nerves in the surgery group and 28 nerves in the medical group. After one year the mean difference was 3.08 (95% CI 2.12 to 4.04) in the surgery group and 2.26 (95% CI 0.21 to 4.31) in the medical group indicating a mean improvement in both. The improvement was greater in the surgery group but the mean difference 0.82 (95% CI -1.34 to 2.98) between the two groups was not significant (Figure 3; Analysis 1.3). In the surgery group 20 out of 31 nerves (65%) had motor improvement compared with 22 out of 31 nerves (71%) in the medical group. The difference was not significant (relative risk 0.91, 95% CI 0.64 to 1.28) (Figure 4; Analysis 1.4).
Figure 4. Forest plot of comparison: Medial epicondylectomy plus oral steroids versus oral steroids alone, outcome: Proportion of ulnar nerves with motor improvement after one year.

<table>
<thead>
<tr>
<th>Study or Subgroup</th>
<th>Treatment Events</th>
<th>Control Events</th>
<th>Weight</th>
<th>Risk Ratio M.H, Fixed, 95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Penmikar 1984</td>
<td>20</td>
<td>22</td>
<td>100.0%</td>
<td>0.91 [0.64, 1.28]</td>
</tr>
<tr>
<td>Total (95% CI)</td>
<td>31</td>
<td>31</td>
<td>100.0%</td>
<td>0.91 [0.64, 1.28]</td>
</tr>
</tbody>
</table>

Heterogeneity: Not applicable
Test for overall effect: Z = 0.54 (P = 0.59)
Secondary outcome measures

1) Improvement in nerve function two years after registration

The trial measured nerve function improvement of the ulnar nerve two years after admission (n = 57 participants with 75 nerves). Results were available for 29 nerves in the surgery group and 28 nerves in the medical group. Improvement was measured as a change score between baseline and end of follow up. Mean differences between the baseline score and the score at the end of two years were compared for the two treatment groups. After two years the mean difference in sensory score was 2.89 (95% CI 0.94 to 4.84) in the surgery group and 2.91 (95% CI 0.73 to 5.09) in the medical group indicating a mean improvement in both. The improvement was slightly greater in the medical group but the mean difference -0.02 (95% CI -2.82 to 2.78) between the two groups was not significant (Figure 5; Analysis 1.5). The mean difference in motor score after two years was 2.79 (95% CI 1.03 to 4.55) in the surgery group and 2.57 (95% CI 0.49 to 4.65) in the medical group indicating a mean improvement in both. The improvement was greater in the surgery group but the mean difference 0.22 (95% CI -2.39 to 2.83) between the two groups was not significant (Figure 6; Analysis 1.6).

(2) Change in nerve pain and in nerve tenderness one year after registration

Pannikar et al evaluated nerve pain and tenderness one year after registration using the scale as defined by Pearson (Pearson 1982). At the end of one year nerve pain and tenderness had disappeared in both groups. Ebenezer et al reported no new nerve pain or tenderness between the first and second year.

(3) Changes in quality of life

The trial did not evaluate changes in quality of life.

(4) Occurrence of adverse events
The trial did not report any adverse events or reasons of loss to follow-up. Contacting the authors did not yield additional information.

**Longitudinal epineurotomy and nerve decompression plus oral corticosteroids versus oral corticosteroids alone for participants with neuritis of less than six months duration (Boucher 1999)**

**Primary outcome measures**

1) Improvement in sensory nerve function one year after registration

The trial did not report results one year after treatment.

2) Improvement in motor nerve function one year after registration

The trial did not report results one year after treatment.

**Secondary outcome measures**

1) Improvement in nerve function two years after registration

The trial compared oral prednisolone plus epineurotomy and external nerve decompression (surgery group) with oral prednisolone alone (medical group) in participants with neuritis of less than six months duration \( (n = 31 \text{ participants}) \). The trial measured sensory nerve function improvement of ulnar, median and posterior tibial nerves and motor nerve function improvement of ulnar, median and common peroneal nerves two years after treatment. Results were available for 46 nerves in the surgery group and 47 nerves in the medical group. Improvement was measured as a change score between baseline and end of follow up and converted into median improvement. For example, a median improvement of 25% means that 50% of the data had greater than 25% improvement and 50% of the data had less than 25% improvement. Sensory testing was done with five graded Semmes-Weinstein monofilaments \( (50 \text{ mg}, 200 \text{ mg}, 2 \text{ g}, 4 \text{ g}, 10 \text{ g}) \) and according to a protocol described by Pearson \( (Pearson 1982) \). Sensory improvement was defined as a positive difference between the final and initial score. Outcomes were expressed as median improvement, meaning that 50% of the data had greater improvement than this value and 50% of the data had less improvement than this median. In the surgery group median sensory improvement was 25% compared to 20% median improvement in the medical group. The difference was not significant at a 5% level (Tukey box plot test) \( (Analysis 2.2) \). No numbers, test values or 95% confidence interval values were given. Contacting the author revealed that original data were no longer available.

2) Change in nerve pain and in nerve tenderness one year after registration

Boucher et al evaluated nerve pain and tenderness two years after registration using the scale as defined by Pearson \( (Pearson 1982) \). In the surgery group median nerve pain relief was 11% compared to 0% in the medical group. The difference was significant at a 5% level (Tukey box plot test) \( (Analysis 2.3) \). No numbers, test values or 95% confidence interval values were given. Contacting the author revealed that original data were no longer available.

3) Changes in quality of life

The trial did not evaluate changes in quality of life.

4) Occurrence of adverse events

One participant was excluded from the study due to haemorrhage, but it was unclear if it was caused by the intervention.

**DISCUSSION**

**Summary of main results**

Decompressive surgery is frequently used in the management of nerve damage in leprosy, but evidence from randomised controlled trials for the effect of decompressive surgery is scarce.

**Overall completeness and applicability and quality of the evidence**

Two randomised controlled trials were available for this review. One trial compared the added benefit of medial epicondylectomy over corticosteroids for participants with ulnar neuritis of less than six months duration \( (Ebenezer 1996; Pannikar 1984) \). The other trial compared the added benefit of longitudinal epineurotomy over corticosteroids for participants with ulnar, median, common peroneal or posterior tibial nerve involvement of less than six months duration \( (Boucher 1999) \). The interventions and outcomes were too heterogeneous to be combined in a meta-analysis. The numbers of participants included in the trials were small and did not allow for subgroup analysis. The variability between studies and the limitations in study design and sample size made it difficult to draw any robust conclusions.

None of the trials found a significant difference in improved nerve function between surgery and medical groups after a follow-up of one or two years. This result may have been biased by the
selection criteria used for inclusion of patients and nerves. Only a small proportion (about 10%) may benefit from decompressive surgery and show improvement after surgery (Naafs 2008). The other nerves need no decompression. By taking all nerves together, results may be diluted and the conclusion clouded. The two trials had some drawbacks. One major drawback of both trials was that they sometimes used more than one nerve from individual patients in the analyses thereby considering the outcomes from each nerve independently. The trial of Pannikar and Ebenezer included 18 patients with ulnar nerve damage at both sides (bilateral involvement). The right side was allocated to the group drawn by random selection and the left side was allocated to the other group. The final results reflect the outcomes of all nerves. No separate analysis was done using only one independent outcome from each patient. Original data were not available. The trial of Boucher included 31 patients with 93 nerves in total. It is unclear how many nerves each patient contributed. The final results reflect the outcomes of all nerves. No separate analysis was done using only one independent outcome from each patient. Original data were not available. Results from these studies should be treated with considerable caution, because results from a patient contributing outcomes from more than one nerve will be treated, in the analysis, as having more weight as a patient contributing only one nerve.

Other limitations of the study of Pannikar were that randomisation was done by alternation, which is considered an inadequate randomisation procedure. With regard to loss to follow-up, 23% of the participants were lost to follow-up after one year and 32% after two years. No reasons for these losses were reported and no intention-to-treat analysis was performed. The randomisation procedure and loss to follow-up (6%) were considered adequate in the study of Boucher. Outcomes were expressed as median improvement. No numbers or original data were available to calculate mean differences or relative risks making comparison and interpretation of the results difficult. Subgroup analyses showed no difference in median improvement between operated and non-operated nerves with respect to type of leprosy (lepromatous or non-lepromatous), type of antibacterial drug therapy (mono or multi), type of nerve function impairment (motor or sensory), and duration of neuritis (0 to 3 months or 3 to 6 months). There were significant differences for pain relief and severity of the neuritis before surgery. Operated nerves had higher median pain relief compared to non-operated nerves. In the group with considerable loss of nerve function the operated nerves had higher median improvement compared to non-operated nerves. The occurrence of adverse effects was not adequately reported in the trials. One study (Boucher 1999) excluded a participant with haemorrhage during the course of the trial, but it was unclear whether this was due to the intervention. The literature reviewing decompressive surgery in leprosy often does not take adverse effects into account, but stresses the importance of having adequate techniques and instruments and competent surgeons to prevent unfavourable outcomes (Bernardin 1997; Bourtrel 1992; Richard 2004). Complications of decompressive surgery in general may be painful scars, wound problems, haematoma, infection and damage to nerves, arteries or tendons (Malaviya 2004a; Scholten 2007; Thoma 2004). None of the trials included quality of life measures or cost-effectiveness calculations which could be useful indicators of the effectiveness of interventions.

Potential biases in the review process

The search process was elaborate and to our knowledge no other randomised controlled trials were available for this review.

Agreements and disagreements with other studies or reviews

Many published and unpublished non-randomised studies have examined the effect of decompressive surgery for treating nerve damage in leprosy. While the two RCTs give insufficient evidence in favour of decompressive surgery in addition to steroid treatment, most non-randomised studies report beneficial effects of decompressive surgery. Relief of nerve pain and tenderness is the most frequently and consistently reported benefit. Nerve function improvement is frequently reported, but the response to surgery seems to depend on several factors, such as severity and duration of neuritis before surgery, the type of leprosy, the nerve involved and the surgical technique used. Nerves which are partially damaged, have neuritis of less than six months duration and are associated with multibacillary (MB) leprosy show better results (Chaise 2004; Kazen 1996; Malaviya 2004b; Palande 1980; Pandya 1983). Studies examining the effects of surgery reported sensory improvement varying from about 38% to 97% and motor improvement varying from about 26% to 63% (Anita 1976; Bernardin 1997; Brandsma 1983; Chaise 1982; Chaise 1985; Chaise 1987; Husain 1997; Husain 1998; Kamar 1982; Malaviya 1982; Palande 1973; Pandya 1978; Ramaroses 1995). Comparison of these studies is difficult due to differences in surgical techniques used, duration and severity of neuritis, type of leprosy, follow up time, and outcome measures.

Several non-randomised studies compared operated versus non-operated nerves. One study evaluated nerve function in nine individuals with neuritis of less than six months duration. Three patients underwent ulnar nerve decompression, three patients received corticosteroid therapy for ulnar neuritis and three patients underwent median nerve decompression. The study found an average nerve function improvement of 35% for ulnar nerve decompression (n = 3), 32% for steroid treatment of eight weeks (n = 3) and 18%, for median nerve decompression (n = 3) six months after surgery or start of treatment (Shah 1986). Three studies examined surgery alone versus surgery plus steroids. One study compared medial epicondylectomy alone (n = 7) with medial epicondylectomy plus steroids (n = 7) given two weeks.
postoperatively for ulnar neuritis of less than one month duration. After a 5-month follow-up motor improvement was not better in the group receiving additional steroids (Oommen 1979). Another study compared neurolysis (n = 21) with neurolysis in combination with perineural corticosteroid injections (n = 18) for ulnar neuritis of less than six months duration. The injections were administered around the thickened nerve after surgery and two and three weeks later. One year after surgery the mean difference between final and initial nerve function score was 14 for the surgery only group and 21 for the surgery plus steroid group (Dandapat 1991). The third study compared decompressive surgery alone (n = 59) with surgery plus steroids (n = 25) given for 3 to 4 months for sensory impairment of the posterior tibial nerve of varying duration. Satisfactory recovery of nerves with duration of anaesthesia of less than six months was 61% in the surgery group and 83% in the surgery plus steroids group four weeks after surgery (Rao 1989).

One study compared operated nerves with contralateral non-operated nerves. Prior to surgery all participants had received three months of steroid treatment. The most affected nerves underwent surgical decompression and were compared with the contralateral non-operated nerves one year or more after surgery. Of the more than 100 nerve decompressions four operated nerves had decreased nerve function after one year of follow up. The other operated nerves had unchanged or improved nerve function one year after surgery. It is unclear how many of the contralateral non-operated nerves improved or deteriorated (Droogenbroeck 1977). After losses to follow-up, another study compared operated nerves (n = 195) of 95 patients with non-operated nerves of 96 patients, matched for type of leprosy, age and duration of sensory loss but not randomised, on changes in sensation. Participants in whom no improvement of sensory nerve function was found after a standard steroid treatment (40 mg prednisolone daily for three weeks after which the dosage was reduced by 5 mg per week) were included in the study. Between 27% and 66% of the nerves had definite improvement two years after surgery compared to 7% of the non-operated nerves (Theuvenet 2006). Improvement was more likely if the sensory loss had been present for a shorter time. Studies from Carayon et al favour surgery plus medical treatment above medical treatment alone (Carayon 1985a; Carayon 1985b; Carayon 1993). Corticosteroids are the cornerstone of management in acute nerve damage in leprosy, are recommended by the WHO and are widely available. But corticosteroids have some shortcomings. The effects of corticosteroids in the long-term remain uncertain and a considerable proportion of people treated for nerve damage do not benefit from corticosteroid treatment. Long-term therapy may cause serious adverse effects, such as peptic ulcer, cataract or psychosis. Spontaneous improvement or recovery of nerve function in untreated or placebo treated individuals has been reported and needs more investigation. The limitations of corticosteroids urge the need to find alternative therapeutic approaches (Van Veen 2007). Surgery alone as therapy for treating neuritis is not recommended, but there is discussion about whether the combination of surgery and medical treatment (e.g. steroids) will give better results than medical treatment alone and there is a call for appropriate trials examining this question (Bourrel 1992; Malaviya 2004b; Richard 2004).

**Authors’ Conclusions**

**Implications for practice**

Evidence from the two randomised controlled trials is insufficient to draw robust conclusions about the effect of decompressive surgery for treating nerve damage in leprosy. Two trials examining the added benefit of surgery over steroids for neuritis of less than six months duration did not show significantly better outcomes with steroids plus surgery than steroids alone in the long-term. Adverse effects of decompressive surgery for treating nerve damage in leprosy are not well-documented.

**Implications for research**

There is a need to identify factors which will predict a favourable response to decompressive surgery or groups of patients or nerves that will be likely to benefit from surgery. Future randomised controlled trials should be well-designed to establish the usefulness and effectiveness of the combination of decompressive surgery and medical treatment compared to medical treatment alone. New trials should pay more attention to non-clinical aspects, such as costs and impact on quality of life, because these are highly relevant indicators for both policy makers and participants.

**Acknowledgements**

We would like to thank Dr P Bourrel, Dr M Ebenezer, Dr J Millan and Dr B Naafs for providing additional information and the Cochrane Neuromuscular Disease Group for advice and help.
References to studies included in this review

Boucher 1999 [published data only]

Ebenezer 1996 [published data only]

Pannikar 1984 [published data only]

References to studies excluded from this review

Carayon 1985a [published data only]

Carayon 1985b [published data only]

Carayon 1993 [published data only]

Dandapat 1991 [published data only]

Droogenbroeck 1977 [published data only]

Oommen 1999 [published data only]

Rao 1989 [published data only]

Additional references

Antia 1976

Becc-Bleumink 1990

Bernardin 1997

Bourrel 1992

Brandsma 1981

Brandsma 1983

Britton 1998

Chaise 1982

Chaise 1985

Chaise 1987

Chaise 2004

Croft 1999
Decompressive surgery for treating nerve damage in leprosy (Review)

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Croft 2000

Heijnders 2004

Husain 1997

Husain 1998

ILEP 2001

ILEP 2002

Job 1989

Juni 2001

Kazen 1996

Kumar 1982

Leekassa 2004

Lockwood 1993

Lockwood 2000

Lockwood 2005

Malaviya 1982

Malaviya 2004a

Malaviya 2004b
Malaviya GN. Shall we continue with nerve trunk decompression in leprosy?. Indian Journal of Leprosy 2004;76(4):331–42.

Marlowe 2004

Meima 2008

Naafs 1996

Naafs 2008
Letter to authors 2008 August. Personal correspondence.

Nicholls 2003

Palande 1973

Palande 1980

Pandya 1978

Pandya 1983

Pearson 1982

Rafferty 2005
Decompressive surgery for treating nerve damage in leprosy (Review)

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## Characteristics of Studies

### Characteristics of included studies [ordered by study ID]

#### Boucher 1999

| Methods | Randomised, parallel group trial  
|         | Randomisation by a computer random number table  
<table>
<thead>
<tr>
<th></th>
<th>Blinding not possible</th>
</tr>
</thead>
</table>
| Participants | 31 leprosy patients with nerve deficit < 6 months duration  
|         | Unit of randomisation: ulnar, median, common peroneal or posterior tibial nerve.  
|         | Unit of analysis: nerve  
|         | Nerves randomised: unclear  
|         | Nerves analysed: 93 (a: 47, b: 46) |
| Interventions | (a) Prednisone start at 40 mg/day for 15 days and thereafter gradually tapered with 5 mg/15 or 30 days until 6 months completed (total 3450 mg)  
|         | (b) Same intervention plus external nerve decompression and a simple, longitudinal epineurotomy |
| Outcomes | Change in:  
|         | (1) Sensory score after 2 years  
|         | (2) Voluntary muscle testing (VMT) score after 2 years  
|         | (3) Nerve pain after 2 years |
| Notes | Single centre  
|         | Conducted in Senegal |

#### Risk of bias

<table>
<thead>
<tr>
<th>Item</th>
<th>Authors' judgement</th>
<th>Description</th>
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</thead>
<tbody>
<tr>
<td>Allocation concealment?</td>
<td>Yes</td>
<td>A - Adequate</td>
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</tbody>
</table>

#### Ebenezer 1996

| Methods | Randomised, parallel group trial  
|         | Randomisation by alternation  
<table>
<thead>
<tr>
<th></th>
<th>Blinding not possible</th>
</tr>
</thead>
</table>
| Participants | 57 leprosy patients with ulnar neuritis < 6 months duration  
|         | Unit of randomisation: person  
|         | Unit of analysis: ulnar nerve  
|         | Persons randomised: 57 with 75 ulnar nerves (18 bilateral cases)  
|         | Nerves analysed: 57 of 39 persons (a: 28, b: 29) |
| Interventions | (a) Prednisolone 30 mg/day for 1 week, reducing the daily dose by 5 mg every week for 6 weeks (total 735 mg)  
(b) Same intervention plus external nerve decompression and a simple, subperiosteal medial epicondylec-tomy |
|----------------|----------------------------------------------------------------------------------------------------------|
| Outcomes | Change in:  
(1) Sensory score after 2 years  
(2) VMT score after 2 years |
| Notes | Single centre  
Conducted in India  
Follow-up study of Pannikar et al |

**Risk of bias**

<table>
<thead>
<tr>
<th>Item</th>
<th>Authors' judgement</th>
<th>Description</th>
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</thead>
<tbody>
<tr>
<td>Allocation concealment?</td>
<td>No</td>
<td>C - Inadequate</td>
</tr>
</tbody>
</table>

**Pannikar 1984**

| Methods | Randomised, parallel group trial  
Randomisation by alternation  
Blinding not possible |
|----------------|----------------------------------------------------------------------------------------------------------|
| Participants | 57 leprosy patients with complaints suggestive of ulnar nerve dysfunction < 24 weeks duration  
Unit of randomisation: person  
Unit of analysis: ulnar nerve  
Persons randomised: 57 with 75 ulnar nerves (18 bilateral cases)  
Nerves analysed: 62 of 44 persons (a: 31, b: 31) |
| Interventions | a) Prednisolone 30 mg/day for 1 week, reducing the daily dose by 5 mg every week for 6 weeks (total 735 mg)  
(b) Same intervention plus external nerve decompression and a simple, subperiosteal medial epicondylec-tomy |
| Outcomes | Change in:  
(1) Sensory score after 1 year  
(2) VMT score after 1 year  
(3) Nerve pain and tenderness after 1 year  
(4) Stretch test |
| Notes | Single centre  
Conducted in India |
### Risk of bias

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<th>Description</th>
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### Characteristics of excluded studies  [ordered by study ID]

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<td>Carayon 1985a</td>
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<tr>
<td>Carayon 1985b</td>
<td>Unclear randomisation procedure</td>
</tr>
<tr>
<td>Carayon 1993</td>
<td>Unclear randomisation procedure</td>
</tr>
<tr>
<td>Dandapat 1991</td>
<td>Unclear randomisation procedure</td>
</tr>
<tr>
<td>Droogenbroeck 1977</td>
<td>No randomisation procedure</td>
</tr>
<tr>
<td>Oommen 1979</td>
<td>No randomisation procedure</td>
</tr>
<tr>
<td>Rao 1989</td>
<td>No randomisation procedure</td>
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</table>
## DATA AND ANALYSES

### Comparison 1. Medial epicondylectomy plus oral steroids versus oral steroids alone

<table>
<thead>
<tr>
<th>Outcome or subgroup title</th>
<th>No. of studies</th>
<th>No. of participants</th>
<th>Statistical method</th>
<th>Effect size</th>
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</thead>
<tbody>
<tr>
<td>Change in sensory score after one year</td>
<td>1</td>
<td>57</td>
<td>Mean Difference (IV, Fixed, 95% CI)</td>
<td>0.08 [-2.45, 2.61]</td>
</tr>
<tr>
<td>Proportion of ulnar nerves with sensory improvement after one year</td>
<td>1</td>
<td>62</td>
<td>Risk Ratio (M-H, Fixed, 95% CI)</td>
<td>1.13 [0.71, 1.77]</td>
</tr>
<tr>
<td>Change in motor score after one year</td>
<td>1</td>
<td>57</td>
<td>Mean Difference (IV, Fixed, 95% CI)</td>
<td>0.82 [-1.34, 2.98]</td>
</tr>
<tr>
<td>Proportion of ulnar nerves with motor improvement after one year</td>
<td>1</td>
<td>62</td>
<td>Risk Ratio (M-H, Fixed, 95% CI)</td>
<td>0.91 [0.64, 1.28]</td>
</tr>
<tr>
<td>Change in sensory score after two years</td>
<td>1</td>
<td>57</td>
<td>Mean Difference (IV, Fixed, 95% CI)</td>
<td>-0.02 [-2.82, 2.78]</td>
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<tr>
<td>Change in motor score after two years</td>
<td>1</td>
<td>57</td>
<td>Mean Difference (IV, Fixed, 95% CI)</td>
<td>0.22 [-2.39, 2.83]</td>
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### Comparison 2. Longitudinal epineurotomy plus oral steroids versus oral steroids alone

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<th>No. of participants</th>
<th>Statistical method</th>
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<tr>
<td>Median sensory improvement after two years</td>
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<td>Other data</td>
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<tr>
<td>Median motor improvement after two years</td>
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<td>Other data</td>
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<tr>
<td>Median improvement in nerve pain and tenderness after two years</td>
<td>1</td>
<td>Other data</td>
<td>No numeric data</td>
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</table>
Analysis 1.1. Comparison 1 Medial epicondylectomy plus oral steroids versus oral steroids alone, Outcome 1 Change in sensory score after one year.

Review: Decompressive surgery for treating nerve damage in leprosy

Comparison: 1 Medial epicondylectomy plus oral steroids versus oral steroids alone

Outcome: 1 Change in sensory score after one year

<table>
<thead>
<tr>
<th>Study or subgroup</th>
<th>Treatment</th>
<th>Control</th>
<th>Mean Difference</th>
<th>Weight</th>
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<tr>
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<td>Mean(SD)</td>
<td>N</td>
<td>Mean(SD)</td>
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<tr>
<td>Ebenezer 1996</td>
<td>29</td>
<td>2.08 (4.74)</td>
<td>28</td>
<td>2 (5)</td>
<td>100.0 %</td>
</tr>
<tr>
<td>Total (95% CI)</td>
<td>29</td>
<td>28</td>
<td>100.0 %</td>
<td>0.08 [ -2.45, 2.61 ]</td>
<td></td>
</tr>
</tbody>
</table>

Heterogeneity: not applicable

Test for overall effect: Z = 0.06 (P = 0.95)

Analysis 1.2. Comparison 1 Medial epicondylectomy plus oral steroids versus oral steroids alone, Outcome 2 Proportion of ulnar nerves with sensory improvement after one year.

Review: Decompressive surgery for treating nerve damage in leprosy

Comparison: 1 Medial epicondylectomy plus oral steroids versus oral steroids alone

Outcome: 2 Proportion of ulnar nerves with sensory improvement after one year

<table>
<thead>
<tr>
<th>Study or subgroup</th>
<th>Treatment</th>
<th>Control</th>
<th>Risk Ratio</th>
<th>Weight</th>
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<tr>
<td></td>
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<td>Pannikar 1984</td>
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<td>16/31</td>
<td>1.13 [ 0.71, 1.77 ]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total (95% CI)</td>
<td>31</td>
<td>31</td>
<td>100.0 %</td>
<td>1.13 [ 0.71, 1.77 ]</td>
<td></td>
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</table>

Total events: 18 (Treatment), 16 (Control)

Heterogeneity: not applicable

Test for overall effect: Z = 0.51 (P = 0.61)
Analysis 1.3. Comparison 1  Medial epicondylectomy plus oral steroids versus oral steroids alone, Outcome 3 Change in motor score after one year.

Review: Decompressive surgery for treating nerve damage in leprosy

Comparison: 1 Medial epicondylectomy plus oral steroids versus oral steroids alone

Outcome: 3 Change in motor score after one year

<table>
<thead>
<tr>
<th>Study or subgroup</th>
<th>Treatment</th>
<th>N</th>
<th>Mean (SD)</th>
<th>Control</th>
<th>N</th>
<th>Mean (SD)</th>
<th>Mean Difference</th>
<th>Weight</th>
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</tr>
</thead>
<tbody>
<tr>
<td>Ebenezer 1996</td>
<td>29</td>
<td></td>
<td>3.08 (2.53)</td>
<td>28</td>
<td></td>
<td>2.26 (5.29)</td>
<td>-0.82 [ -1.34, 2.98 ]</td>
<td>100.0 %</td>
<td></td>
</tr>
<tr>
<td>Total (95% CI)</td>
<td>29</td>
<td></td>
<td>3.08 (2.53)</td>
<td>28</td>
<td></td>
<td>2.26 (5.29)</td>
<td>-0.82 [ -1.34, 2.98 ]</td>
<td>100.0 %</td>
<td></td>
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</tbody>
</table>

Heterogeneity: not applicable

Test for overall effect: Z = 0.74 (P = 0.46)

Analysis 1.4. Comparison 1 Medial epicondylectomy plus oral steroids versus oral steroids alone, Outcome 4 Proportion of ulnar nerves with motor improvement after one year.

Review: Decompressive surgery for treating nerve damage in leprosy

Comparison: 1 Medial epicondylectomy plus oral steroids versus oral steroids alone

Outcome: 4 Proportion of ulnar nerves with motor improvement after one year

<table>
<thead>
<tr>
<th>Study or subgroup</th>
<th>Treatment</th>
<th>n/N</th>
<th>Control</th>
<th>n/N</th>
<th>Risk Ratio</th>
<th>Weight</th>
<th>Risk Ratio</th>
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</thead>
<tbody>
<tr>
<td>Pannikar 1984</td>
<td>20/31</td>
<td></td>
<td>22/31</td>
<td></td>
<td>0.91 [ 0.64, 1.28 ]</td>
<td>100.0 %</td>
<td>0.91 [ 0.64, 1.28 ]</td>
</tr>
<tr>
<td>Total (95% CI)</td>
<td>31</td>
<td></td>
<td>31</td>
<td></td>
<td></td>
<td>100.0 %</td>
<td></td>
</tr>
</tbody>
</table>

Total events: 20 (Treatment), 22 (Control)
Heterogeneity: not applicable
Test for overall effect: Z = 0.54 (P = 0.59)
Analysis 1.5. Comparison 1 Medial epicondylectomy plus oral steroids versus oral steroids alone, Outcome 5 Change in sensory score after two years.

Review: Decompressive surgery for treating nerve damage in leprosy

Comparison: 1 Medial epicondylectomy plus oral steroids versus oral steroids alone

Outcome: 5 Change in sensory score after two years

<table>
<thead>
<tr>
<th>Study or subgroup</th>
<th>Treatment</th>
<th>Control</th>
<th>Mean Difference</th>
<th>Weight</th>
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<td>Mean(SD)</td>
<td>Mean(SD)</td>
<td>IV,Fixed,95% CI</td>
<td>IV,Fixed,95% CI</td>
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<tr>
<td>Ebenezer 1996</td>
<td>29</td>
<td>2.89 (5.13)</td>
<td>28</td>
<td>2.91 (5.62)</td>
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<tr>
<td><strong>Total (95% CI)</strong></td>
<td><strong>29</strong></td>
<td><strong>28</strong></td>
<td>100.0 %</td>
<td>-0.02 [-2.82, 2.78]</td>
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Heterogeneity: not applicable

Test for overall effect: Z = 0.01 (P = 0.99)

Analysis 1.6. Comparison 1 Medial epicondylectomy plus oral steroids versus oral steroids alone, Outcome 6 Change in motor score after two years.

Review: Decompressive surgery for treating nerve damage in leprosy

Comparison: 1 Medial epicondylectomy plus oral steroids versus oral steroids alone

Outcome: 6 Change in motor score after two years

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<th>Control</th>
<th>Mean Difference</th>
<th>Weight</th>
<th>Mean Difference</th>
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<td>Mean(SD)</td>
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<td>IV,Fixed,95% CI</td>
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<tr>
<td>Ebenezer 1996</td>
<td>29</td>
<td>2.79 (4.63)</td>
<td>28</td>
<td>2.57 (5.37)</td>
<td>100.0 %</td>
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<tr>
<td><strong>Total (95% CI)</strong></td>
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<td><strong>28</strong></td>
<td>100.0 %</td>
<td>0.22 [-2.39, 2.83]</td>
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Heterogeneity: not applicable

Test for overall effect: Z = 0.17 (P = 0.87)
Median sensory improvement after two years

<table>
<thead>
<tr>
<th>Boucher 1999</th>
<th>25% median improvement</th>
<th>20% median improvement</th>
<th>Tukey box plot test</th>
<th>No significant difference at 5% level</th>
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</table>

Median motor improvement after two years

<table>
<thead>
<tr>
<th>Boucher 1999</th>
<th></th>
<th></th>
<th></th>
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</thead>
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Median improvement in nerve pain and tenderness after two years

<table>
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<th>Boucher 1999</th>
<th>11% median improvement</th>
<th>0% median improvement</th>
<th>Tukey box plot test</th>
<th>Significant difference at 5% level</th>
</tr>
</thead>
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**APPENDICES**

**Appendix 1. OVID MEDLINE search strategy**

1 leprosy.mp. or exp Leprosy/
2 hansen disease.mp.
3 1 or 2
4 exp Decompression/ or decompression$.mp.
5 neurolysis.mp.
6 epicondylectomy.mp.
7 epineurotomy.mp.
8 or/4-7
9 neuritis.mp. or Neuritis/
10 nerve damage.mp.
11 peripheral nervous system diseases.mp. or exp Peripheral Nervous System Diseases/
12 nerve loss.mp.
13 Peripheral Nerves/
14 neuropath$.mp.
15 nerve function impairment.mp.
16 nerve problem.mp.
17 nerve involvement.mp.
18 (nerve pain or neuralgia).mp. or Neuralgia/
19 or/9-18 (165847)
20 randomized controlled trial.pt.
21 controlled clinical trial.pt.
22 randomized controlled trials/
23 random allocation/
24 double-blind method/
25 single-blind method/
26 or/20-25
27 animals/ not humans/
28 26 not 27
29 clinical trial.pt.
30 exp clinical trial/
31 (clin$ adj25 trial$).ti,ab.
32 ((sing$ or doubl$ or tripl$ or trebl$) adj25 (blind$ or mask$)).ti,ab.
Appendix 2. OVID EMBASE search strategy

1 leprosy.mp. or exp Leprosy/
2 hansen disease.mp.
3 1 or 2
4 exp Decompression/ or decompression$.mp.
5 neurolysis.mp.
6 epicondylectomy.mp.
7 epineurotomy.mp.
8 4 or 7
9 neuritis.mp. or Neuritis/
10 nerve damage.mp.
11 peripheral nervous system diseases.mp. or exp Peripheral Nervous System Diseases/
12 nerve loss.mp.
13 Peripheral Nerves/
14 neuropath$.mp.
15 nerve function impairment.mp.
16 nerve problem.mp.
17 nerve involvement.mp.
18 (nerve pain or neuralgia).mp. or Neuralgia/
19 4 or 18
20 Randomized Controlled Trial/
21 Clinical Trial/
22 Multicenter Study/
23 Controlled Study/
24 Crossover Procedure/
25 Double Blind Procedure/
26 Single Blind Procedure/
27 exp RANDOMIZATION/
28 Major Clinical Study/
29 PLACEBO/
30 Meta Analysis/
31 phase 2 clinical trial/ or phase 3 clinical trial/ or phase 4 clinical trial/
32 (clin$ adj25 trial$).tw.
33 ((sing$ or doubl$ or tripl$ or trebl$) adj25 (blind$ or mask$)).tw.
Appendix 3. OVID AMED search strategy

1 leprosy.mp. or exp Leprosy/
2 hansen disease.mp.
3 1 or 2
4 exp Decompression/ or decompression$.mp.
5 neurolysis.mp.
6 epicondylectomy.mp.
7 epineurotomy.mp.
8 or/4-7
9 neuritis.mp. or Neuritis/
10 nerve damage.mp.
11 peripheral nervous system diseases.mp. or exp Peripheral Nervous System Diseases/
12 nerve loss.mp.
13 Peripheral Nerves/
14 neuropath$.mp.
15 nerve function impairment.mp.
16 nerve problem.mp.
17 nerve involvement.mp.
18 (nerve pain or neuralgia).mp. or Neuralgia/
19 or/9-18
20 Randomized controlled trials/
21 Random allocation/
22 Double blind method/
23 Single-Blind Method/
24 exp Clinical Trial/
25 (clin$ adj25 trial$).tw.
26 ((sing$ or doubl$ or treb$ or trip$) adj25 (blind$ or mask$ or dummy)).tw.
27 placebo/
28 placebo$.tw.
29 random$.tw.
30 research design/
31 Prospective Studies/
32 cross over studies.mp.et.
33 meta analysis/
34 (meta?analys$ or systematic review$).tw.
35 control$.tw.
36 (multicenter or multicentre).tw.
Appendix 4. LILACS search strategy

((Pt randomized controlled trial OR Pt controlled clinical trial OR Mh randomized controlled trials OR Mh random allocation OR Mh double-blind method OR Mh single-blind method) AND NOT (Ct animal AND NOT (Ct human and Ct animal)))

OR (Pt clinical trial OR Ex E05.318.760.535$ OR (Tw clin$ AND (Tw trial$ OR Tw ensa$ OR Tw estud$ OR Tw experim$ OR Tw investiga$)) OR ((Tw singl$ OR Tw simple$ OR Tw dobl$ OR Tw doble$ OR Tw duplo$ OR Tw trebl$ OR Tw trip$) AND (Tw blind$ OR Tw cego$ OR Tw mask$ OR Tw mascar$)) OR Mh placebo OR Tw placebo$ OR (Tw random$ OR Tw randon$ OR Tw casual$ OR Tw acaso$ OR Tw azar OR Tw aleator$) OR Mh research design) AND NOT (Ct animal AND NOT (Ct human and Ct animal))

OR (Ct comparative study OR Ex E05.337$ OR Mh follow-up studies OR Mh prospective studies OR Tw control$ OR Tw prospectiv$ OR Tw volunt$ OR Tw volunteer$) AND NOT (Ct animal AND NOT (Ct human and Ct animal))

AND (Mh leprosy OR Tw hansen$ disease OR Tw leprosy) AND (Mh Decompression OR Tw decompression OR Tw neurolysis OR Tw epicondylectomy OR Tw epineurotomy) AND (Tw neuritis OR Mh neuritis OR Tw neuropath$ OR Tw neuropath$ OR Tw nerve damage OR Tw nerve involvement OR Tw nerve loss OR Tw nerve function impairment OR Tw nerve problem OR Mh peripheral nerve$ OR Tw peripheral nervous system disease$ OR Mh peripheral nervous system diseases OR Tw nerve pain OR Tw neuralgia OR Mh neuralgia)

Appendix 5. CINAHL search strategy

S40. S39 and S21
S39. S38 and S26 and S21
S38. S37 or S36 or S35 or S34 or S33 or S32 or S31 or S30 or S29 or S28 or S27
S37. nerve pain
S36. (neuralgia) or (MH "Neuralgia")
S35. nerve involvement
S34. nerve problem
S33. nerve function impairment
S32. neuropath*
S31. (Peripheral Nerves) or (MH "Peripheral Nerves")
S30. nerve loss
S29. (peripheral nervous system diseases) or (MH "Peripheral Nervous System Diseases")
S28. nerve damage
S27. (neuritis) or (MH "Neuritis")
S26. S25 or S24 or S23 or S22
S25. epineurotomy
S24. epicondylectomy
S23. neurolysis
S22. (Decompression) or (MH "Decompression, Surgical")
S21. S20 or S19
S20. hansen disease
S19. (leprosy) or (MH "Leprosy")
S18. S17 or S16 or S15 or S14 or S13 or S12 or S11 or S10 or S9 or S8 or S7 or S6 or S5 or S4 or S3 or S2 or S1
S17. TI random* or AB random*
S16. (TI (cross?over or placebo* or control* or factorial or sham? or dummy) OR AB (cross?over or placebo* or control* or factorial or sham? or dummy))
S15. (TI clin* or intervention* or compar* or experiment* or prevent* or therapeutic or AB clin* or intervention* or compar* or experiment* or prevent* or therapeutic) and (TI (trial*) or AB (trial*))
S14. (TI meta?analys* or systematic review*) or (AB meta?analys* or systematic review*)
S13. (TI single* or doubl* or tripl* or trebl*) or AB (single* or doubl* or tripl* or trebl*) and (TI blind* or mask*) or AB (blind* or mask*)
S12. ABAB design*
S11. PT clinical trial or PT systematic review
S10. (MH “Factorial Design”)
S9. (MH “Concurrent Prospective Studies”) or (MH “Prospective Studies”)
S8. (MH “Meta Analysis”)
S7. (MH “Solomon Four-Group Design”) or (MH “Static Group Comparison”)
S6. (MH “Quasi-Experimental Studies”)
S5. (MH “Placebos”)
S4. (MH “Double-Blind Studies”) or (MH “Triple-Blind Studies”)
S3. (MH “Clinical Trials+”)
S2. (MH “Crossover Design”)
S1. (MH “Random Assignment”) or (MH “Random Sample”) or (MH “Simple Random Sample”) or (MH “Stratified Random Sample”) or (MH “Systematic Random Sample”)

HISTORY
Protocol first published: Issue 1, 2008
Review first published: Issue 1, 2009

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<th>Date</th>
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<td>14 November 2007</td>
<td>New citation required and conclusions have changed</td>
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CONTRIBUTIONS OF AUTHORS
Link with editorial base and co-ordinate contributions from co-authors (NvV).
Draft protocol (NvV with input from all).
Run search (NvV).
Identify relevant titles and abstracts from searches (NvV, JHR).
Obtain copies of trials (NvV).
Selection of trials (NvV, JHR).
Extract data from trials (NvV, JHR).
Enter data into RevMan (NvV).
Carry out analysis (NvV, JHR).
Interpret data (NvV, TS, WT, JHR).
Draft final review (NvV with input from all).
Update review (NvV, JHR).
DECLARATIONS OF INTEREST

None declared.

SOURCES OF SUPPORT

Internal sources

- The Netherlands Leprosy Relief, Netherlands.

External sources

- No sources of support supplied