Content and outcome of usual primary care for back pain: a systematic review

Simon Somerville, Elaine Hay, Martyn Lewis, Julie Barber, Danielle van der Windt, Jonathan Hill and Gail Sowden

ABSTRACT
Background
Most patients seeking help for back pain are managed in primary care.

Aim
To describe the content and outcome of ‘usual care’ for low back pain in primary care trials.

Design of study
A systematic review of randomised controlled trials published since 1998.

Setting
Primary care.

Method
Randomised controlled trials of back pain in adults were scrutinised to obtain data on treatment and outcome measures in groups receiving usual primary care. A narrative review of the resulting heterogeneous data was undertaken.

Results
Thirty-three papers were identified for analysis. Overall the exact nature of the treatment received in the ‘usual’ primary care group was poorly recorded. Medication was frequently used, and there were suggestions that levels of opioid prescription were higher than might be expected from clinical guidelines. Requesting of plain-film X-rays occurred more often than recommended. There was very little information to suggest that doctors were promoting physical activity for patients with back pain. Disability scores (Roland–Morris Disability Questionnaire) and pain scores improved over time for patients with acute or subacute back pain, but not for those with chronic pain.

Conclusion
Treatment received by patients with back pain was varied and often not in line with back-pain guidelines, particularly with respect to opioid prescription and X-ray investigation. The content of the ‘usual care’ arm in trials is crucial to interpreting the outcome of studies, but was poorly described in the papers reviewed. Future trials should more fully describe the ‘usual care’ arm.

Keywords
back pain; family practice; general practice; primary health care.

INTRODUCTION
Low back pain is a common condition that is frequently seen in primary care. In the UK there are between 3 and 7 million GP consultations for back pain in a year. Fifty-two million working days are lost each year and, at any one time, 6% of the working population will have had at least one day off work in the last 4 weeks due to back pain.7 The annual cumulative consultation rate of patients presenting to their GP with back pain is 6.4%.2

Despite evidence-based guidelines for the management of low back pain,24 chronic or recurrent back pain is common and it continues to pose considerable challenges and frustrations for patients and practitioners. Attempts have been made to address this situation using educational programmes for GPs to promote the adoption of guidelines, but these have failed to show significant changes in everyday clinical practice.24 One reason for this may be that the nature of routine clinical practice in primary care has not been adequately considered when producing guidelines.

‘Usual care’ is often used as the control or comparator group in randomised controlled trials. It
is important to know the content of these control treatments in order to interpret the effect sizes published in trials. The current study is a systematic review with the following specific objectives:

- to describe the treatments contained within ‘usual care’ approaches to low back pain, as reported in primary care-based randomised controlled trials published between 1994 and 1997 (a period when many clinical guidelines were published); and
- to describe the clinical course of patients randomised to the ‘usual care’ approaches outlined in the first objective.

**METHOD**

**Inclusion criteria for studies to review**

Types of studies. Randomised controlled trials published in any language from 1998 only were included. This date range was chosen as it follows the publication date of many national guidelines on the care of low back pain in primary care worldwide, and hence reflects current clinical practice.

Types of participants. Participants aged 18 years or over with non-specific low back pain of any duration, who had been treated in primary care, were included. Low back pain was defined as pain in the back located below the scapulae and above the natal cleft of the buttocks, with or without pain radiating to the leg.

Studies involving specific pathological processes or surgical techniques were excluded.

Types of treatments. A randomised controlled trial was included for review if at least one treatment group involved usual care of low back pain in primary care. Any interventions used to provide this usual care were recorded.

Types of outcome measures. For the study’s first objective of describing usual care approaches to treat low back pain, relevant outcome measures were consultation rates, advice, medication usage (prescribed to the patient or purchased by the patient without prescription), investigations, and referrals. For the second objective of describing the clinical course of low back pain in those receiving usual care, information was gathered on pain scores (Visual Analogue Scale, modified Von Korff Pain Scale, Aberdeen Back Pain Scale, and the Extended Aberdeen Spine Pain Scale), overall satisfaction, back-specific functional scores (Roland–Morris Disability Questionnaire [RMDQ]), and work absence.

Search strategy for identification of studies

Relevant studies meeting the inclusion criteria were identified by a computer search of the electronic databases MEDLINE (1966–2007), EMBASE (1966–2007), CINAHL (1982–2007), AMED, DARE, and ISI Web of Science. The search strategy looked for terms within the subject heading, title, and abstract, and included the following terms: random, clinical trials, low back pain, back pain, backache, physicians-family, family practice, general practice, family medicine, primary care, and primary medical care (Appendix 1). The search was augmented by examination of the Cochrane Central Register of Controlled Trials and references given in the identified randomised controlled trials and appropriate reviews.

**Methods of review**

Study selection. The principal researcher performed the initial database search using the search strategy. The number of potential studies was reduced by the removal of duplicates. Studies were excluded if it was immediately apparent from their abstract that they did not meet the inclusion criteria. Trials in the remaining group were then assessed by examining the full paper.

Methodological quality assessment. One researcher scored all papers; each paper was also independently assessed by other researchers. Disagreements about the methodological score were resolved by discussion.

The Cochrane Collaboration Back Review Group publishes guidelines on systematic reviews. It recommended an 11-point scoring system for assessing methodological quality, including items relating to internal and external validity. Studies with a score of ≥6 out of 11 are judged to be of high quality, as they are above the arbitrary cut-off point of 50%.

Subsequently, it has been suggested that the list of assessment criteria can be reduced to just the five points relating to internal validity, which are: (1) concealment of treatment allocation; (2) blinding of patients; (3) blinding of outcome assessor; (4) intention-to-treat analysis; (5) and an acceptable drop-out rate. In this review a study was judged to be of high quality if it achieved a score of ≥3 out of 5.

There was disagreement between reviewers about quality scores of six out of 33 papers. In every case disagreements were resolved by the two researchers involved, without the need to call on a third reviewer to adjudicate.

**How this fits in**

Despite back pain being a common problem in primary care, it is poorly described and its treatment appears at odds with guideline recommendations. A better understanding of primary care is needed to facilitate the incorporation of evidence-based medicine into routine consultations.
Abstracts examined (n = 164)

Papers excluded (n = 110)

Full papers examined (n = 54)

Studies excluded (n = 21): not RCT, not primary care, surgical procedures, or involving specific diseases

RCTs included in review (n = 33)

References identified by literature search (n = 253)

Duplicates removed (n = 89)

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**Figure 1. Systematic review flowchart.**

Data extraction. This was performed by the lead researcher and other independent reviewers. One key aspect of the review was to determine whether usual care had been adequately described in terms of types of treatments that health professionals had at their discretion to use ('allowed'), and what treatments actually took place ('actual'). Each reviewer was instructed to make their own judgement as to whether allowed and actual usual care had been adequately described. Disagreements were resolved by discussion.

A standardised form was used by each reviewer to record their findings. Information gathered was collated using a Microsoft Access database.

Data analysis. Due to the sparseness and heterogeneity of the data, it was appropriate to perform a qualitative assessment rather than a meta-analysis. Also, the purpose of this study was to describe and statistically summarise treatment and outcome measures relating to the usual care arm of trials, rather than to compare effect sizes.

Frequencies were presented of specific elements of care, for example, consultation rates, medication usage (prescribed or purchased over the counter), and referrals. Mean scores with standard deviations (SDs) were extracted from each paper (if stated) and presented for pain (Visual Analogue Scale, modified Von Korff Pain Scale, Aberdeen Back Pain Scale) and RMDQ score. Measures of pain and satisfaction were all converted to a 0–10-point scale (a higher score indicated greater pain or greater satisfaction respectively) to facilitate interpretation and comparison across studies.

Duration of back pain in each study was classified pragmatically as acute (0–6 weeks), subacute (6–12 weeks), chronic (more than 12 weeks), or mixed (any duration included in study). It should be acknowledged that such classification may not reflect clinical practice, where back pain often fluctuates with time and recurrent acute attacks may be indistinguishable from relapses of a chronic problem.

**RESULTS**

**Study selection**

The initial search yielded 164 potentially relevant studies. After examining the full text of 54 papers, 33 were included in the review. This process is detailed in Figure 1.

**Study characteristics**

Study characteristics are presented in detail in Appendix 2. Type of care given or intended to be given in the usual care control group is described, in addition to the outcome of the experimental intervention compared with usual care.

Of the 33 studies identified, eight investigated patients with acute back pain, 13,19,20,23,29,32,35,39,66,31,37 scored 3,1,12,17,19-21,23,27,30,32-41,16 scored 3,1,14-16,19,23,25,28,31,33,35,36,38,39,43 and one scored 5 out of a maximum score of 5. In 16 out of 33 studies it was not possible to tell if the outcome assessor was blinded, 3,12-17,23,27,29,30,34,37,41,43 and in only three of 33 studies were the patients blinded to the nature of the treatment group. In two studies it was not stated if the treatment allocation was concealed, 3,40 and in three it was not possible to tell if the analysis was by intention to treat. All papers reported the drop-out rates.

**Methodological quality**

In the 33 studies, one study scored 1,24 two scored 2,27,30 13 scored 3,17,18,20,23,29,32,37,40-42 16 scored 4,2,14-16,19,23-25,28-29,30,32-40,42 and one scored 5 out of a maximum score of 5. In 16 out of 33 studies it was not possible to tell if the outcome assessor was blinded, 3,12-17,23,27,29,30,34,37,41,43 and in only three of 33 studies were the patients blinded to the nature of the treatment group. In two studies it was not stated if the treatment allocation was concealed, 3,40 and in three it was not possible to tell if the analysis was by intention to treat. All papers reported the drop-out rates.

**Content of usual care approaches**

Adequacy of description of usual care. Of 33 studies, 11 had no description of either the allowed or actual treatment in the control group. Seven had a description of both, respectively to facilitate interpretation and comparison across studies.

Duration of back pain in each study was classified pragmatically as acute (0–6 weeks), subacute (6–12 weeks), chronic (more than 12 weeks), or mixed (any duration included in study). It should be acknowledged that such classification may not reflect clinical practice, where back pain often fluctuates with time and recurrent acute attacks may be indistinguishable from relapses of a chronic problem.

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described just the actual treatment,\textsuperscript{2,22,25,33,43} and nine just the allowed treatment.\textsuperscript{12,13,16,19,20,31,36,37,39}

The 10 most commonly recorded aspects of usual care are presented in Table 1. Of these 10 common aspects, a mean of 3.1 items (SD 2.2) were reported in the studies included in the review. Appendix 3 gives more information about aspects of consultations and demonstrates the wide variation in the detail of recording of usual care.

Consultation rates with GP. Due to the heterogeneous ways in which data about consultations were collected, it was not possible to perform a comprehensive quantitative analysis. Overall, 17 of 33 papers reported data on consultation rates. The following qualitative data were found:

- **Acute back pain.** Four of eight studies described consultation rates per patient, one found a rate of 3.9 (no SD stated) consultations in 1 year,\textsuperscript{22} and another 3.3 (SD 1.6) in 8 weeks.\textsuperscript{15} Another found that 60% of patients presented to their GP once, 31% twice, and 9% more than twice in the course of a year.\textsuperscript{13} A fourth states that consultation rates were similar for patients in the experimental and control arms, although figures are not given.\textsuperscript{39}

- **Subacute back pain.** Four of 11 studies provide data on consultations for subacute back pain. One found 62.5% of patients presented to their GP in the first 3 months of the trial, and 51.1% in the second 3 months.\textsuperscript{33} A second study reported that 21% consulted once, 3% twice, and 5% three or more times in the first 6 months, and 16%, 3%, and 5% respectively at 9 months.\textsuperscript{15} Mean number of visits to the GP in 1 year was six in a further study.\textsuperscript{21} Jellema et al found that 24% of patients were seeking additional help from their GP in the first 6 weeks, and 28% in the remainder of the year of the study.\textsuperscript{18}

- **Chronic back pain.** Two of six studies reported data on consultations for chronic back pain. One reported a mean of 13.2 (SD: 5) visits in one year,\textsuperscript{31} and another reported a mean of 1.2 at baseline for the previous year and 1.7 in the year of the study.\textsuperscript{25}

- **Mixed-duration back pain.** Of eight papers, one found the mean number of visits to be 3.6,\textsuperscript{22} and one found 4.8 visits for mixed-duration back pain.\textsuperscript{22} Thomas et al reported that 40% of patients presented to the GP in the first 12 weeks of their study.\textsuperscript{43}

Medication

Only 12 out of 33 studies gave information about medication taken by patients for their back pain.\textsuperscript{21,31,33,35,36,41,43}

Many patients were taking medication at baseline, with figures reported of 69%,\textsuperscript{20} 80%,\textsuperscript{24} 90%,\textsuperscript{43} and 69% (prescribed) and 73% (over the counter).\textsuperscript{21}

Two papers gave information about opioid prescription: 34% of patients in a US study,\textsuperscript{38} and 18.7% in the UK\textsuperscript{39} were prescribed them. Unfortunately, there was no information about the strength of the opiates or whether combination drugs were prescribed.

Work absence

Information about absence from work was quite heterogeneous due to the way in which authors recorded this information and differences in the type of usual care; some studies only recruited patients who were absent from work because of back pain.\textsuperscript{33,35} It is important to remember this when comparing data about sick notes from different countries. The different procedures for obtaining a sick notes in different countries may give a false interpretation of the data. As mentioned in the paper Dutch GPs do not give sick notes, British GPs do.

**Acute back pain.** One study reported that 81% of patients had no certified sick leave, fewer than 9% had 10 or more days of certified sick leave in a year, and overall there was a mean of 3 days’ sick leave.\textsuperscript{13} Another reported a mean 2.2 days (SD: 3.4) of sick leave since randomisation at 8 weeks’ follow-up.\textsuperscript{15}

**Subacute back pain.** Patients reported a mean of 14 days of sick leave at baseline,\textsuperscript{21} and 15 in the 3 months before commencement of the study.\textsuperscript{15}

**Chronic back pain.** Varied information was found related to work absence for chronic back pain. Two studies reported that 50%,\textsuperscript{21} and 57.3%\textsuperscript{17} of patients were off sick at baseline, whereas in another study the patients only had 3 days off sick at baseline.\textsuperscript{25}

<table>
<thead>
<tr>
<th>Table 1. Ten most commonly recorded aspects of usual care.</th>
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</thead>
<tbody>
<tr>
<td>Consultation activity record</td>
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<tr>
<td>-----------------------------</td>
</tr>
<tr>
<td>Radiology used</td>
</tr>
<tr>
<td>Other investigation ordered</td>
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<tr>
<td>Medication used (prescribed or over the counter)</td>
</tr>
<tr>
<td>Sick note issued</td>
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<tr>
<td>Written advice given</td>
</tr>
<tr>
<td>Activity level recorded</td>
</tr>
<tr>
<td>Exercise advice recorded</td>
</tr>
<tr>
<td>Seen by physiotherapist</td>
</tr>
<tr>
<td>Hospital referral made</td>
</tr>
<tr>
<td>Consultation rate details</td>
</tr>
</tbody>
</table>

See Appendix 3 for reference details.
Activity levels and exercise advice
Three studies gave details on the activity levels of patients before they presented to their GP and three reported the advice given to patients about exercise during the trial.

Physiotherapy. For acute back pain, 7% and 8.6% of patients were referred to physiotherapy. For subacute back pain, 18%, 31%, and 92.9% of patients receiving usual care were referred to physiotherapy at baseline or in the first 6 weeks. For mixed-duration back pain 49% were referred within the first 12 weeks. In the case of chronic back pain it was not possible to compare data in a meaningful way, due to heterogeneity as discussed in relation to work absence.

Radiology. Use of plain film X-rays in the management of back pain varied, particularly between countries. The percentages of patients having this investigation were 19% (US), 53.6% (Canada), 7% (UK), and 4% (UK).

Clinical course of patients randomised to usual care

Functional disability (RMDQ score). The data shown in Figures 2a and 2b suggest that the speed at which RMDQ scores improve is inversely related to the duration of back pain: that is, the longer the duration of pain, the slower the rate of improvement. In acute and subacute pain there was a definite trend for improvement. The single study that did not follow this trend involved usual care provided by university-affiliated clinics and the emergency department of a hospital in an inner-city setting in the US; as such, these patients may not be representative of patients normally managed in primary care.

Pain score. For acute and subacute pain there was a general trend for pain scores to improve with time, as observed on the 0–10–point pain scale used to rate the studies. In two studies acute pain mean pain scores improved by 5.8 and 2.0 between baseline and their endpoints respectively. For subacute pain, mean pain scores improved by 1.2, 5.0, 2.0, 1.0, and 3.9 respectively.

For patients with chronic pain, the pain score changed little over the course of the study, with changes in mean pain scores of 0.9, –0.8, 0.6 and –1.8 respectively (a negative value meaning reduction of pain).

Satisfaction. For acute pain, mean satisfaction scores at study endpoint were 7.2 (treatment), 7.0 (explanation), and 4.1 (overall), according to the 0–10–point satisfaction scale used to rate the studies. In subacute pain, endpoint satisfaction scores were 4.7 and 4.1. In one study on chronic back pain the mean endpoint satisfaction score was 4.4. A study on mixed-duration back pain expressed this differently, with the number of patients who were very satisfied or satisfied at 85% following their initial consultation, and at 70% after 6 weeks (Figure 2c).

DISCUSSION

Methodological quality
That many studies did not have blinding of patients or outcome assessors is not unusual. Often blinding
is not possible, or even sensible, due to the pragmatic nature of the trial design, the interventions used, or ethical constraints.

Studies were presented in different ways, which made quality assessment and data extraction difficult to carry out. Future randomised controlled trials on back pain should be presented in line with Consolidated Standards of Reporting Trials (CONSORT) recommendations.27,32,34

Overall, most papers reviewed were of an acceptable quality according to the study’s quality scale. A small number of trials failed to describe concealment of allocation to treatment,35,40 or intention-to-treat analysis.27,30,34

Direct comparison between studies was difficult due to the differing ways in which data were recorded. For example, pain experienced by patients was assessed using Visual Analogue Scale, modified Von Korff Pain Scale, Aberdeen Back Pain Scale, and the Extended Aberdeen Spine Pain Scale. Nevertheless, the patterns of changes in pain were similar to those for disability, with patients with acute and subacute back pain showing larger improvements than participants with chronic back pain.

**Description of usual care**

Overall, usual care was poorly described in these studies. Historically, there has been a tendency to allow care in the control group to be according to the normal practice of the care provider, with little or no recording of what actually took place. This contrasts with the intervention treatment, against which usual care is compared, which is often described in great detail.

It is clear that healthcare providers vary widely in their preferred practices, due to many factors including their experience and the expectations of the population in which the studies are set. As such, control groups in separate studies reviewed here may receive a range of different treatments. This makes comparison between studies difficult.

Although the practice of comparing a new treatment with the prevailing current practice is well accepted, interpretation of such studies is problematic when the usual care groups vary so widely and are not adequately described. This is particularly pertinent given that a healthcare provider may act differently purely by knowing their practice is under scrutiny.

Attempts to standardise usual care have shown little effect in changing clinician behaviour.3 It is important to accept that usual care is heterogeneous, and this needs to be taken into account when interpreting the results of studies. The present authors suggest that a minimum dataset about usual care is recorded to improve understanding of the treatments received in the usual care group (Table 1 and Appendix 2). This includes information about radiology usage, other investigations ordered, medications used (prescribed or over-the-counter), sick notes issued, use of written materials, recording of activity levels, details of exercise advice given, consultation rates, physiotherapy undertaken, and hospital referrals.5

**Content of usual care**

It is difficult to draw any firm conclusions about how frequently primary care consultation is allowed or what actually happens in the usual care approaches. Marked differences existed in the control groups due to the country involved, variation in the study designs, and methods of recording consultation activity; only half of studies reported data and the results showed no pattern.

Similarly, the overall recording of medication use was quite poor. This is disappointing in that medication use has significant financial costs for the patient and the health economy, and is associated with a not-insignificant number of side-effects.

This review found that many patients take over-the-counter medication for their back pain, and GPs often provided additional medication. Data were limited, but there was a suggestion that levels of opioid prescription were quite high,3,15 given the message in back-pain management guidelines that suggest using weak opioids as ‘add-on’ treatments when paracetamol or non-steroidal anti-inflammatory drugs provide insufficient pain relief.3,46 Compound analgesics are not recommended as they do not allow separate titration of the constituent medications, and there are concerns about dependence with strong opioids.3,45

Unfortunately, in the studies in this review,
information was not recorded about the strength of opiates or whether combination products were used. It may be that the concerns about strong and combination opioids are overemphasised, particularly when the majority of patients will recover from their back pain in a few weeks. A GP faced with a patient who is in pain, despite taking simple painkillers, may well decide that it is appropriate to prescribe these drugs in the short term, but with close monitoring for side-effects and over-usage. It is important to record more detailed information about the types of drugs used to treat back pain, and their effectiveness and safety.

It is generally accepted that plain-film X-rays are not helpful in the management of most patients with back pain and represent an avoidable source of radiation. This review found that the use of X-rays varied widely between countries, from 4% of cases in the UK, to 53.6% in Canada. Reasons for requesting investigations are complex and involve many factors including patient and doctor beliefs and knowledge, country-specific healthcare systems, and patient satisfaction.

The tension between guideline advice and actual clinical behaviour is illustrated by a survey of GPs that found that they would almost always or sometimes refer for X-ray in 70% of patients with non-recurrent backache of less than a month’s duration: 88% per cent of requests were to reassure the patient, and 78% to reassure the doctor. This finding is not confined to low back pain consultations, and it has been shown that perceived patient pressure, rather than clinical need, is the reason behind a significant minority of investigations. Clearly, it is desirable to reduce patient exposure to radiation, but to achieve this goal a guideline must be of practical use to doctors in primary care. The guideline must include adequate explanation of the rationale behind each recommendation, address the needs for reassurance on the part of the patient and doctor, and be adaptable to local resources (for example, it is no use advocating magnetic resonance imaging instead of plain-film X-rays if it is not readily available).

The clear messages from trials and guidelines are that patients with low back pain should continue with normal activities, even if uncomfortable, and stay at or return to work. The papers included in this review did not, in general, report on this important aspect of patient care. One possible reason for this might be that it is not the normal practice of GPs to record this information. Also GPs may not feel able to give specific advice about exercise, due either to lack of training or fear of litigation should the patient develop a problem related to exercising. GPs are well placed to give exercise advice to patients, and there is a need to address these issues to encourage and support people who would benefit from increasing their activity.

**Course of symptoms**

The relatively small number of studies involved makes it difficult to undertake a meaningful data synthesis and arrive at firm conclusions. However, Figures 2a and 2b suggest that the speed at which the RMDQ score changed over time depended on the duration of the back disorder. For acute pain there was a trend for RMDQ scores to fall rapidly over the first few weeks (Figure 2a), and this improvement was maintained throughout the study period. For chronic pain sufferers the outlook was much less promising, and RMDQ scores did not tend to improve with time.

Usual care groups in the studies in this review were given a range of treatments but, despite this, it appears that the effectiveness of usual care (if measured as improvement of RMDQ score) may be related more to the duration of the episode than to the type of care given.

Usual care is variable in content and effectiveness. That it is also poorly described compounds this problem and makes comparison between studies even more difficult. As such, it is important that usual care is not regarded as a uniform control against which interventions are compared without attempting to understand what it consists of. This heterogeneity may undermine the fundamental basis for calculating sample sizes in future trials. Agreement is needed as to what information should be recorded in future trials, and conformity to CONSORT needs to be improved.

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North Staffordshire Primary Care Research Consortium

**Competing interests**

The authors have declared that there are none

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48. Danish Institute for Health Technology Assessment. Low back pain. Frequency, management and prevention from an HTA perspective. Copenhagen: Danish Health Technology Assessment, 1999 [In Danish].
### Appendix 1. Search strategy for identification of studies.

Relevant studies meeting the inclusion criteria were identified by:

1. A computer search of MEDLINE (1966–2007), EMBASE (1966–2007), CINAHL (1982–2007), AMED, DARE, and ISI Web of Science. The following search strategies were used. They were restricted to adults, human participants, and publication since 1998:

   - **MEDLINE**
     
     (random$ OR clinical trial$ [pt, sh, ti, ab]) AND (low back pain OR back pain or backache [sh, ti, ab]) AND (physicians-family OR family pract$ OR general pract$ OR family medicine OR primary health care OR primary medical care [sh, ti, ab])

   - **EMBASE**
     
     (random$ OR clinical trial$ [pt, sh, ti, ab]) AND (low back pain OR back pain OR backache OR lumbago [sh, ti, ab]) AND (physicians-family OR family pract$ OR general pract$ OR family medicine OR primary health care OR primary medical care [sh, ti, ab])

   - **CINAHL**
     
     (random$ OR controlled clinical trial$ [pt, sh, ti, ab]) AND (low back pain OR back pain or backache ) AND (low back pain OR back pain or backache [sh, ti, ab]) AND (physicians-family OR family pract$ OR general pract$ OR family medicine OR primary health care OR primary medical care [sh, ti, ab])

2. Examination of the Cochrane Central Register of Controlled Trials.

3. Examination of references given in the identified randomised controlled trials and appropriate reviews.
### Participants: Baseline mean Outcome of usual intervention

<table>
<thead>
<tr>
<th>Study (first author)</th>
<th>Setting</th>
<th>Duration of back pain</th>
<th>Participants: n, mean age in years, % female</th>
<th>Baseline mean RMĐQ score (SD)</th>
<th>Intervention</th>
<th>Control</th>
<th>Outcome of usual care mean (SD) unless stated</th>
<th>Outcome of intervention in comparison with usual care</th>
<th>Methodological quality score (0–5, with 5 being best)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Brealey**</td>
<td>UK general practice</td>
<td>Mixed</td>
<td>338, 42.5, 53</td>
<td>9 (3.9)</td>
<td>Physical treatments; exercise</td>
<td>Best care</td>
<td>RMĐQ: base 9 (3.9), 3 months 6.8 (3.3), 12 months 6.1 (0.3) versus exercise group. Pain (MVK): base 60.5 (17.6), 3 months 49.3 (1.6), 12 months 48.4 (1.7) versus exercise group</td>
<td>Compared with best care — exercise: small benefit at 3 months, nil at 12 months. Manipulation: small to moderate benefit at 3 months and small benefit at 12 months</td>
<td>3</td>
</tr>
<tr>
<td>Burton**</td>
<td>UK 5 general practices and 1 osteopathy practice</td>
<td>Acute</td>
<td>79, 44.7, 48</td>
<td>9.7 (4.6)</td>
<td>Back book (biopsychosocial) or handy-hints pamphlet (biomedical)</td>
<td>Usual care + pamphlet</td>
<td>RMĐQ: base 9.7 (4.6), 12 months –4.5 (-), Pain (VAS 0–100) worst/best: base 68.7 (18.9)/50.8 (27.8) 12 months 56.3 (18.7)/10.6 (17.8)</td>
<td>Improved beliefs about low back pain; reduced fear-avoidance and RMĐQ score; no effect on pain</td>
<td>5</td>
</tr>
<tr>
<td>Cherkin**</td>
<td>US primary care</td>
<td>Acute</td>
<td>66, 40.1, 42</td>
<td>11.7 (5.4)</td>
<td>Physical therapy; chiropractic booklet</td>
<td>Usual care + booklet</td>
<td>RMĐQ: base 11.7 (1.3), 4 weeks 4.9 (1.1), 12 weeks 4.3 (1.2)</td>
<td>Physical and chiropractic therapy improves satisfaction, but nothing else</td>
<td>4</td>
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<tr>
<td>Curtis**</td>
<td>US primary care</td>
<td>Acute</td>
<td>143, 42.7, 57</td>
<td>15.6 (5.4)</td>
<td>Manual therapy given by GP</td>
<td>Enhanced care patient-centred approach, careful examination, including palpation and functional assessment, use of national guidelines and patient handouts regarding activities of daily living and exercise. Option to use standard prescription, bed rest, trigger-point injections referral to others</td>
<td>RMĐQ at baseline 15.8 (5.4); improvement in RMĐQ from randomisation: base 5.7, 2 weeks 8.7 (8.0), 4 weeks 9.8 (7.3), 8 weeks 11.1 (7.9), Pain (0–10): 4.6 (2.1). Satisfaction (1–5, 1 = best); endpoint 2.1 (1.1)</td>
<td>GP giving limited manual therapy offers very modest benefit compared with enhanced care alone. However, only 43% of patients in manual treatment group given full treatment course by the GP</td>
<td>4</td>
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<tr>
<td>Damush***</td>
<td>US inner-city health centres and emergency rooms</td>
<td>Subacute</td>
<td>106, 45.5, 75</td>
<td>13.9 (6.8)</td>
<td>Self-management programme for patients on low income with acute low back pain: long-term outcomes (1 year)</td>
<td>Usual care referral to occupational health, physical therapy, or a neurologist; analgesics, back exercise advice sheets</td>
<td>RMĐQ: base 13.9 (6.8), 12 months 11.3 (8.1), Satisfaction (higher score better): base 24.3 (5.7), 12 months 25.4 (5.4)</td>
<td>At 12 months patients who participated in the self-management programme showed significantly better RMĐQ scores, mental functioning, self-efficacy to manage acute low back pain, time in physical activity, and reduced fear of movement/re-injury</td>
<td>4</td>
</tr>
<tr>
<td>Day*</td>
<td>UK primary care</td>
<td>Mixed</td>
<td>1138, 41.3, 54.3</td>
<td>–</td>
<td>Education of GPs to promote use of guidelines on low back pain</td>
<td>Usual care</td>
<td>Patients receiving the following (%): X-ray 13.7, sick note 19.2, opiates/muscle relaxant 18.7, referral to secondary care 2.3, referral to physiotherapy 13.8</td>
<td>No difference between study and control patients in terms of proportion referred to X-ray, given a sick certificate, prescribed opioids or muscle relaxants, or referred to secondary care. Only 21% of patients had evidence that they were given advice about avoiding bed rest; 0.6% in both arms recommended to take bed rest. Suggested that GPs know about guidelines but feel pressurised by patient expectations to behave differently</td>
<td>4</td>
</tr>
<tr>
<td>Molsø Hagen**</td>
<td>Norway spine clinic</td>
<td>Chronic</td>
<td>220, 41.1, 48</td>
<td>–</td>
<td>Light mobilisation</td>
<td>Usual care</td>
<td>19.6 days of compensated sick leave in 2 years</td>
<td>Brief examination, information giving, reassurance and encouragement to resume physical activity reduces long-term sick leave in the first year. Those returning to work mainly do so in the first year</td>
<td>3</td>
</tr>
</tbody>
</table>

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**Continued...**
## Appendix 2. Study characteristics (continued).

<table>
<thead>
<tr>
<th>Study (first author)</th>
<th>Setting</th>
<th>Duration of back pain</th>
<th>Participants: mean age in years, % female</th>
<th>Baseline mean RMDQ score (SD)</th>
<th>Intervention</th>
<th>Control</th>
<th>Outcome of usual care: mean (SD) unless stated</th>
<th>Outcome of intervention in comparison with usual care</th>
<th>Methodological quality score (0–5, with 5 being best)</th>
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</thead>
<tbody>
<tr>
<td>Hagen *</td>
<td>Norway spine clinic</td>
<td>Subacute</td>
<td>457, 41, 48</td>
<td>12.2 (6); median 13 IQR (8–16)</td>
<td>Assessment and modification of psychological factors</td>
<td>Usual care as per Dutch College of GP guidelines</td>
<td>RMDQ: base 12.2 (6.0); base 13 (8–16), 6 weeks 4 (1–10), 13 weeks 2 (0–5), 26 weeks 1 (0–3), 1 year 1 (0–4). Pain 0–10; base 5 (5–6), 6 weeks 2 (0–6), 13 weeks 1 (0–3), 26 weeks 0 (0–2), 1 year 0 (0–2)</td>
<td>No difference in outcome measures versus usual care</td>
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<td>Jellem a *</td>
<td>The Netherlands primary care</td>
<td>Subacute</td>
<td>171, 42, 47</td>
<td>12.2 (6); median 13 IQR (8–16)</td>
<td>Assessment and modification of psychological factors</td>
<td>Usual care as per Dutch College of GP guidelines</td>
<td>RMDQ: base 12.2 (6.0); base 13 (8–16), 6 weeks 4 (1–10), 13 weeks 2 (0–5), 26 weeks 1 (0–3), 1 year 1 (0–4). Pain 0–10; base 5 (5–6), 6 weeks 2 (0–6), 13 weeks 1 (0–3), 26 weeks 0 (0–2), 1 year 0 (0–2)</td>
<td>No difference in outcome measures versus usual care</td>
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<td>Karjalainen *</td>
<td>Finnish primary care</td>
<td>Subacute</td>
<td>57, 43, 60</td>
<td>12.2 (6); median 13 IQR (8–16)</td>
<td>Assessment and modification of psychological factors</td>
<td>Usual care as per Dutch College of GP guidelines</td>
<td>RMDQ: base 12.2 (6.0); base 13 (8–16), 6 weeks 4 (1–10), 13 weeks 2 (0–5), 26 weeks 1 (0–3), 1 year 1 (0–4). Pain 0–10; base 5 (5–6), 6 weeks 2 (0–6), 13 weeks 1 (0–3), 26 weeks 0 (0–2), 1 year 0 (0–2)</td>
<td>No difference in outcome measures versus usual care</td>
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<tr>
<td>Karjalainen *</td>
<td>Finland primary care</td>
<td>Subacute</td>
<td>57, 43, 60</td>
<td>12.2 (6); median 13 IQR (8–16)</td>
<td>Assessment and modification of psychological factors</td>
<td>Usual care as per Dutch College of GP guidelines</td>
<td>RMDQ: base 12.2 (6.0); base 13 (8–16), 6 weeks 4 (1–10), 13 weeks 2 (0–5), 26 weeks 1 (0–3), 1 year 1 (0–4). Pain 0–10; base 5 (5–6), 6 weeks 2 (0–6), 13 weeks 1 (0–3), 26 weeks 0 (0–2), 1 year 0 (0–2)</td>
<td>No difference in outcome measures versus usual care</td>
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<td>Kendrick *</td>
<td>UK primary care</td>
<td>Subacute</td>
<td>211, 39, 60</td>
<td>8 (4–12)</td>
<td>X-rays for low back pain</td>
<td>Usual care as per Dutch College of GP guidelines</td>
<td>RMDQ: base 8 (4–12), 3 months 3 (1–7), Pain 0–10; base 2 (1–6), 3 months 3 (1–7), Satisfaction (EuroQol): base 20 (17.75–22), 3 months 19 (19–23)</td>
<td>X-rays don't improve care of low back pain suffers, but increase doctors' workload</td>
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<td>Kerry *</td>
<td>UK primary care</td>
<td>Mixed</td>
<td>322, 41.1, 53</td>
<td>10.8 (5.4)</td>
<td>X-rays for first presentation of low back pain</td>
<td>Usual care as per Dutch College of GP guidelines</td>
<td>RMDQ: base 10.8 (5.4), 6 weeks 5.4 (0.3), 1 year 4.2 (0.3). Pain (SF-36 BP): base 4.5 (26), 6 weeks 56 (2), 1 year 65 (2)</td>
<td>Referral for lumbar X-ray doesn't improve pain, function, or disability. There is a small possible improvement in psychological status, but need to balance against radiation risk</td>
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<td>Kovacs *</td>
<td>Mallorca primary care</td>
<td>Chronic</td>
<td>45, 45.6, 66.7</td>
<td>12.17 (-)</td>
<td>Neuroreflexotherapy</td>
<td>Usual care as per Dutch College of GP guidelines</td>
<td>RMDQ: base 12.2 (6.5–18.0), 1-year improvement 2.1 (–1.5 to 6.7), Pain (VAS): base 5.2 (4.1–8), 1-year improvement 1.9 (–1.3 to 2.3)</td>
<td>Neuroreflexotherapy reduced pain, disability, specialist referral, X-ray requests, drug costs, and sick leave</td>
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<td>Licciardone *</td>
<td>US university clinic</td>
<td>Chronic</td>
<td>20, 49, 65</td>
<td>7.3 (5.4)</td>
<td>Osteopathic manipulation in chronic low back pain</td>
<td>Usual care as per Dutch College of GP guidelines</td>
<td>RMDQ: base 7.3 (5.4), Pain (VAS): base 3.1 (2.3), 6 months −3.9 (–), Global satisfaction: base −1.9 (–), 6 months −2.1 (–)</td>
<td>Osteopathic manipulation improved pain, treatment satisfaction, function, and mental health compared with usual care. Sham manipulation improved pain, treatment satisfaction, and function compared with usual care. There were no differences between osteopathic and sham manipulation</td>
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<td>Linton *</td>
<td>Sweden primary care</td>
<td>Chronic</td>
<td>70, 45, 71</td>
<td>–</td>
<td>CBT and different forms of information</td>
<td>Usual care as per Dutch College of GP guidelines</td>
<td>Pain (0–10): base 4.8 (0.45), 1 year 4.0 (0.55)</td>
<td>CBT caused a nine-fold reduction in sick leave and physician and physiotherapy usage</td>
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<td>Meng *</td>
<td>US various clinics in private and hospital sectors</td>
<td>Chronic</td>
<td>24, 70, 62.5</td>
<td>11.8 (5.3)</td>
<td>Acupuncture for chronic low back pain</td>
<td>Usual care as per Dutch College of GP guidelines</td>
<td>RMDQ: base 11.8 (5.3), 6 weeks improved by 0.6 (2.7), Pain (VAS): base 1.7 (1.3), 6 weeks worsened by 0.6 (1.2)</td>
<td>Acupuncture reduced RMDQ scores and was safe</td>
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<tr>
<td>Study</td>
<td>Setting</td>
<td>Participants</td>
<td>Baseline mean RMDQ score (SD)</td>
<td>Intervention</td>
<td>Control</td>
<td>Outcome of usual care: mean (SD) unless stated</td>
<td>Outcome of intervention in comparison with usual care</td>
<td>Methodological quality score (0–5, with 5 being best)</td>
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<tr>
<td>Miller</td>
<td>UK primary care</td>
<td>211, 39, 16</td>
<td>8 (4–12)</td>
<td>Cost effectiveness of X-rays</td>
<td>Usual care</td>
<td>Median (IQR) RMDQ: base 8 (4–12) Pain (VAS, 0–100)</td>
<td>X-rays are only cost-effective when patient satisfaction is high. Other ways of obtaining satisfaction are to be preferred</td>
<td>2</td>
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<td>Kalber, Moffett</td>
<td>UK primary care</td>
<td>98, 42, 56</td>
<td>5.56 (3.34)</td>
<td>Exercise programme</td>
<td>Usual care</td>
<td>RMDQ: base 5.56 (3.94), % of patients with improvement ≥2 points; 6 months 47, 1 year 47.5</td>
<td>High fear-avoiders benefit most from an exercise programme and the effect is maintained at 1 year. Patients who are depressed/distressed gained a short-term benefit</td>
<td>4</td>
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<tr>
<td>Moffett</td>
<td>UK primary care</td>
<td>98, 42, 56</td>
<td>5.56 (3.34)</td>
<td>Exercise programme</td>
<td>Usual care</td>
<td>RMDQ: base 5.56 (3.94), improvement from base; 6 weeks 9.9, 6 months 8.11, 1 year 8.48</td>
<td>Exercise programme reduced pain, disability, and healthcare usage</td>
<td>3</td>
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<tr>
<td>Moore</td>
<td>US HMO clinics</td>
<td>113, 49, 49.6</td>
<td>8.29 (5.8)</td>
<td>Self-care (CBT)</td>
<td>Book on back care</td>
<td>RMDQ: 8.29 (5.8), 3 months 6.55 (6.13), 6 months 6.4 (5.99), 1 year 5.96 (5.8), Pain (VAS –0–100): base 5.2 (1.98), 3 months 4.06 (2.71), 6 months 3.71 (2.28), 1 year 2.98 (1.99)</td>
<td>CBT approach reduced back-related worry, fear-avoidance, pain, and functional problems</td>
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<tr>
<td>Moseley</td>
<td>Australia physiotherapy clinic</td>
<td>28, 398, 54</td>
<td>11.9 (3.2)</td>
<td>Combined physiotherapy and education</td>
<td>Usual care</td>
<td>RMDQ: base 11.9 (3.2), 1-year improvement 4.3, Pain (0–10): base 4.7 (1.5), 1-year improvement 1.4</td>
<td>A combined programme of manual therapy, exercise, and education reduced pain and disability</td>
<td>4</td>
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<tr>
<td>Roberts</td>
<td>UK primary care</td>
<td>28, 29, 2, 32</td>
<td>–</td>
<td>Back-care leaflet</td>
<td>Back-care</td>
<td>–</td>
<td>The leaflet improved posture but had no effect on function</td>
<td>3</td>
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<tr>
<td>Rossignol</td>
<td>Canadian worker compensation scheme</td>
<td>56, 38, 23.2</td>
<td>–</td>
<td>Advice and support for physicians treating low back pain</td>
<td>Usual care</td>
<td>Pain (VAS, 0–100): base 52.4 (20.7), 3 months 10.9 (24.0), 6 months 12.8 (27.0), Satisfaction (0–55, best–worst); base 30.3 (22.6), little change during trial</td>
<td>Advice and support to doctors about management of low back pain reduces patients sick leave, pain, dysfunction, and X-ray usage. It improves exercise levels</td>
<td>4</td>
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<tr>
<td>Schectman</td>
<td>US HMO clinics</td>
<td>590, 45, 54</td>
<td>–</td>
<td>Physician education versus patient educational material versus neither</td>
<td>Usual care</td>
<td>–</td>
<td>Education of physicians improves compliance with low back pain guidelines. Patient educational material has no effect</td>
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<tr>
<td>Seferlis</td>
<td>Sweden hospital clinic</td>
<td>60, 38, 47</td>
<td>–</td>
<td>Manual therapy versus intensive training versus usual care</td>
<td>Usual care</td>
<td>–</td>
<td>Most patients return to work by 1 month regardless of treatment. Satisfaction is improved by manual therapy or intensive training</td>
<td>4</td>
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<tr>
<td>Seferlis</td>
<td>Swedish hospital clinic previous study</td>
<td>60, 38, 47</td>
<td>–</td>
<td>Cost-minimisation analysis of conservative treatment for back pain</td>
<td>Usual care</td>
<td>–</td>
<td>Usual care has lowest cost</td>
<td>4</td>
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<tr>
<td>Skouen</td>
<td>Norway hospital clinic</td>
<td>86, 44, 64</td>
<td>–</td>
<td>Extensive versus light multidisciplinary treatments versus usual care</td>
<td>Usual care</td>
<td>–</td>
<td>Light programme more effective than usual care. Extensive programme no more effective than usual care</td>
<td>3</td>
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<tr>
<td>Staal</td>
<td>The Netherlands occupational health airline</td>
<td>67, 37, 7</td>
<td>13 (4.9)</td>
<td>Graded activity</td>
<td>Usual care</td>
<td>RMDQ: base 13.0 (4.9), improvement frame base: 3 months 4.9 (6.2), 6 months 6.4 (6.6), Pain (0–10): base 6.4 (1.7), improvement frame base: 3 months 2.5 (2.8), 6 months 2.7 (2.8)</td>
<td>Graded activity reduces days of work lost due to low back pain</td>
<td>4</td>
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<tr>
<td>Thomas</td>
<td>UK acupuncture clinics and primary care</td>
<td>241, 44, 58</td>
<td>–</td>
<td>Acupuncture</td>
<td>Usual care</td>
<td>Disability (ODI) baseline 31.4 (14.2), 12 months 19.6 (5.4), 24 months 21.0 (14.2), Pain (SF-36 BP, 0–100): baseline 30.4 (18.0), 12 months 23.3 (22.2), 24 months 59.5 (23.4), Pain (MPQ): baseline 2.7 (1.0), 12 months 1.53 (0.9), 24 months 1.71 (1.1)</td>
<td>Weak evidence that acupuncture has an effect on persistent back pain at 2 months, stronger evidence of a smaller effect at 24 months</td>
<td>3</td>
<td></td>
</tr>
</tbody>
</table>
### Appendix 2. Study characteristics (continued)

<table>
<thead>
<tr>
<th>Study (first author)</th>
<th>Setting</th>
<th>Duration of back pain</th>
<th>Participants: n, mean age in years, % female</th>
<th>Baseline mean RMDQ score (SD)</th>
<th>Intervention</th>
<th>Control</th>
<th>Outcome of usual care: mean (SD) unless stated</th>
<th>Methodological quality score (0–5, with 5 being best)</th>
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<tbody>
<tr>
<td>UnderwoodEE</td>
<td>UK primary care</td>
<td>Acute</td>
<td>40, 41, 45</td>
<td>Back extension exercises</td>
<td>Usual care</td>
<td>Disability (ODI); base 35.6, 1 year 24% patients had a score &lt;20. Pain (VAS); base 50.4, 1 year 18% had score &lt;20</td>
<td>No evidence to show that it is more effective than usual care</td>
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<tr>
<td>Von KorffEE</td>
<td>US primary care</td>
<td>Mixed</td>
<td>126, 50.3, 56</td>
<td>Lay-led self-management</td>
<td>Usual care</td>
<td>RMDQ (approximate); base 9.0, 3 months 7.4, 6 months 7.3, 1 year 7.0. Pain (average); base 5.7 (2.1), 3 months 4.0 (2.1), 6 months 4.1 (0.9), 1 year 3.8 (2.4)</td>
<td>Self-help group reduced worried, produced positive attitudes, and reduced activity limitations</td>
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<tr>
<td>WilliamsEE</td>
<td>UK primary care</td>
<td>Subacute</td>
<td>106, –, –</td>
<td>–</td>
<td>Effectiveness and healthcare costs of a primary care osteopathy service</td>
<td>Usual care</td>
<td>Pain (VAS, 0–100); base 46.3 (22.6), 2 months 6.8 (23.4), 6 months 10.1 (24.1)</td>
<td>Primary care osteopathy improved short-term physical and long-term psychological outcomes at a little extra cost</td>
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</table>

*Media (interquartile range). ALBPS = Aberdeen Low Back Pain Scale (0–100, 0 = best). CBT = cognitive behavioural therapy. HMO = health maintenance organization. IQR = interquartile range. MPQ = McGill Pain Questionnaire (0–5, 0 = best). MVK = modified Von Korff Pain Scale (0–100, 0 = best). ODI = Oswestry Disability Index (0–100, 0 = best). SF-16 BP = SF-16 Bodily Pain (0–100, 0 = best). RMDQ = Roland-Morris Disability Questionnaire (0–24, 0 = best). VAS = Visual Analogue Scale (0–100 or 0–10, best = 0).*
### Appendix 3. Record of activities within the consultation.

<table>
<thead>
<tr>
<th>Study (first author)</th>
<th>Case mix</th>
<th>Radiology figures</th>
<th>Other investigations</th>
<th>Any drug details (prescribed or OTC)</th>
<th>Sick note detail</th>
<th>Written advice recorded</th>
<th>Activity level details</th>
<th>Exercise referral rate</th>
<th>Physiotherapy details</th>
<th>Hospital consultation details</th>
<th>Consultation rate details</th>
<th>Item score (out of 10)</th>
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<tr>
<td>Cherkins</td>
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<td>Y</td>
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<td>Underwood</td>
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Number of studies giving data: 11, 2, 11, 22, 9, 3, 4, 15, 8, 17

% of studies giving data: 33, 6, 33, 67, 27, 9, 12, 45, 24, 52

OTC = over the counter. Y = number of items recorded (out of 10).