

We use cookies to enhance your experience on our website. By clicking 'continue' or by continuing to use our website, you are agreeing to our use of cookies. You can change your cookie settings at any time.

[Continue](#)
[Find out more](#)

Oxford Journals Medicine British Medical Bulletin Volume 94, Issue 1 Pp. 81-144.

British Medical Bulletin

bmb.oxfordjournals.org

Br Med Bull (2010) 94 (1): 81-144. doi: 10.1093/bmb/ldp052

First published online: January 10, 2010

Rating scales for low back pain

Umile Giuseppe Longo[†], Mattia Loppini[†], Luca Denaro[‡], Nicola Maffulli[§] and Vincenzo Denaro[†]

[+](#) Author Affiliations

* Correspondence to: Nicola Maffulli, Centre for Sports and Exercise Medicine, Barts and The London School of Medicine and Dentistry Mile End Hospital, 275 Bancroft Road, London E1 4DG, UK. E-mail: n.maffulli@qmul.ac.uk

Accepted November 24, 2009.

Abstract

Introduction During the past decades several rating scales have been developed to assess the functional status of patients with low back pain.

Methods We performed a search using the keywords 'spine' in combination with 'scoring system', 'scale', 'scores', 'outcome assessment', 'low back pain' and 'clinical evaluation'.

Results Twenty-eight scoring systems are currently available for the evaluation of low back pain. Each of them evaluates low back pain using specific variables. All these scoring systems are presented.

Discussion Although many scoring systems have been used to evaluate the back function, we are still far from a single outcome evaluation system that is reliable, valid and sensitive to clinically relevant changes, taken into account both patients' and physicians' perspective and is short and practical to use.

Conclusion Further studies are required to evaluate the reliability, validity and sensitivity of the low back pain scoring systems used in the common clinical practice.

Key words [low back pain](#) [rating scores](#) [assessment](#)

Introduction

Low back pain (LBP) is a common symptom, affecting more than 80% of the general population in the industrialized world.¹ It is the most frequent cause of disability in people under 45 years of age.² It represents a relevant social and economic problem in developed countries, being the first reason for orthopaedic consultations.³

The development of instruments to measure the outcome of patient with LBP has been the subject of increasing interest. During the past decades, several score systems have been developed to assess the functional status of patients with LBP.⁴⁻⁷ Many generic and disease-specific measures are available for orthopaedic clinical and research practice.⁸ Self-report questionnaires of pain and functional status allow one to evaluate patients before and after a given treatment, and they can be used to detect short-term or long-term clinical changes of symptoms and disabilities.⁹

The aim of this review is to report all the available score systems for the evaluation of LBP and their use in the current orthopaedic practice.

Materials and methods

Oxford Index

About the Index



Show related links

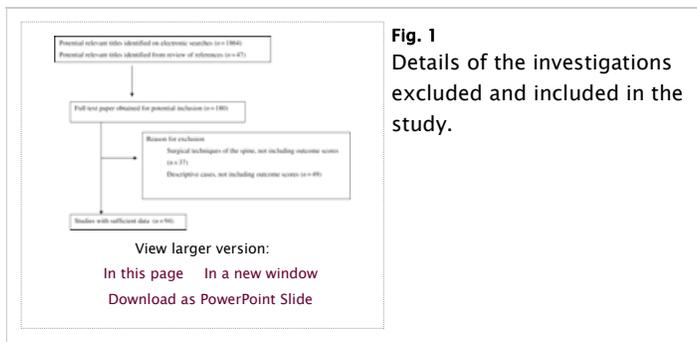
Search across all sources

in Oxford Index



captured 12/17/13

We performed a search using the keywords 'spine' in combination with 'scoring system', 'scale', 'scores', 'outcome assessment', 'low back pain' and 'clinical evaluation', with no limit regarding the year of publication. The following databases were accessed on 20 February 2009: PubMed (<http://www.ncbi.nlm.nih.gov/sites/entrez/>); Ovid (<http://www.ovid.com>); Cochrane Reviews (<http://www.cochrane.org/reviews/>). Given the linguistic capabilities of the research team, we considered publications in English, Spanish and Italian. Two authors (U.G.L. and M.L.) independently read the abstract of each publication identified (if an abstract was available). If no abstract was available, the publication was excluded. In addition, the references section of all the publications identified were studied to ascertain whether other relevant material could be found. The personal collection of scientific material of the senior authors (N.M. and V.D.) was consulted for the same purpose. If deemed relevant, all relevant publications were retrieved. The most relevant material was drawn between the years 1990 and 2007. A large number of publications focusing on surgical techniques of the lumbar spine, not including outcome scores, were not included. The publications thus selected were examined by all authors. After this further selection, 94 publications relevant to the topic at hand were included (Fig. 1).



Analytical description of LBP scoring systems

Roland–Morris disability questionnaire

The Roland–Morris disability questionnaire¹⁰ is constructed by choosing statements from the sickness impact profile (SIP), which is a 136-item health status measure covering a range of aspects of daily living about physical and mental function.^{11,12} The scale consists of 24 yes/no items related specifically to physical functions to specifically assess the disability from LBP (Table 1). The physical functions considered include walking, bending over, sitting, lying down, dressing, sleeping, self-care and daily activities. Patients are asked whether the statements apply to them that day (i.e. the last 24 h). In the scale, one point is given for each item. The RDQ score can be obtained by adding up the number of items checked. The final score ranges from 0 (no disability) to 24 (severe disability). The questionnaire is self-administered by the patient, it can be completed in a maximum of 5 min, and an un-weighted score can be calculated in less than 1 min.

View this table: In this window In a new window	Table 1 The Roland–Morris disability questionnaire.
------------------------------------------------------------------------------------	---------------------------------------------------------------

The original RDQ also contains a six-point pain rating scale in the form of a pain thermometer.¹⁰ However, the authors prefer to use the pain scale of SF-36 instead of scale described in the original article.¹³

Variants of Roland–Morris disability questionnaire

The RDQ-23⁷ (Table 2) is a modified 23-item version of Roland–Morris

disability scale. In this instrument five original items are deleted and replaced with other four items which the authors selected from the SIP. The five deleted items are 'Because of my back, I lie down to rest more often', 'Because of my back, I try to get other people to do things for me', 'My appetite is not very good because of my back', 'Because of my back pain, I get dressed with help from someone else' and 'I sit down for most of the day because of my back'. Each item is scored with 0 or 1 point and the final score is obtained by adding up item scores, ranging from 0 to 23.

View this table:
[In this window](#) [In a new window](#)

Table 2
 The Roland–Morris disability questionnaire 23–item version.

The RDQ–18¹⁴ (Table 3) is a shorter modified version of Roland–Morris disability scale, in which items 2, 15, 17, 19, 20 and 24 are deleted. The other questions are the same of original questions. Item reduction is obtained by measuring the frequency of item endorsement, calculating the inter–item correlations and determining the internal consistency of the questionnaire. The original version scoring scheme is maintained.

View this table:
[In this window](#) [In a new window](#)

Table 3
 The Roland–Morris disability questionnaire 18–item version.

The RDQ–16^{15,16} (Table 4) is a modified 16–item Roland–Morris disability scale designed to measure the limitations in daily living in the past 2 weeks due to back pain. Each item can be answered as follows: 'yes', 'no', 'don't know' or 'not applicable'. For scoring, the number of affirmative answers is divided by the number of questions answered. The final score is expressed by the percentage of items checked with higher scores representing greater limitations.

View this table:
[In this window](#) [In a new window](#)

Table 4
 The modified 16–item Roland–Morris scale.

The RDQ–two time version (RDQ–two)¹⁷ (Table 5) is a modified version of Roland–Morris disability scale produced to assess the LBP over the preceding 4 weeks. Patients have to mark how many days in the previous 4 weeks they had been affected by LBP. Each question has different possible answers and is scored with a 4–week time scale, ranging from 0 to 1 and according to how many days patient is affected: 'not at all' or 'not applicable' are scored with 0 points; '1–7 days' is scored with 0.2 points; '8–14 days' is scored with 0.4 points; '15–21 days' is scored with 0.6 points; '21–27 days' is scored with 0.8 points and 'every day' is scored with 1 point. The number of questions and the domains investigated are the same of the original version. The final score is calculated by dividing the patient total score by the maximum possible score (24) to express the result as a percentage.

View this table:
[In this window](#) [In a new window](#)

Table 5
 The Roland–Morris disability questionnaire two.

The RDQ–7p¹⁸ (Table 6) is a modified version of Roland–Morris disability scale in which a seven point Likert scale is used. This version consists of original wording and original scheme. For scoring, yes/no responses are

labelled as follows: 0 points means 'disagree totally', 3 points means 'not sure' and 6 means 'agree totally'. The final questionnaire score is expressed as percentages of the total possible score with higher scores representing greater disability.

View this table:
[In this window](#) [In a new window](#)

Table 6
 The Roland–Morris disability questionnaire 7p.

The RDQ–12 (Table 7), also named the Maine–Seattle back questionnaire, is a 12–item version of Roland–Morris disability scale derived from the RDQ–23.¹⁹ It is a short self–administered back–specific questionnaire. Like the original scale, the final score is obtained with an unweighted sum of each item score. Thus, the RDQ–12 score can range from 0 (no impairment) to 12 (severe impairment).

View this table:
[In this window](#) [In a new window](#)

Table 7
 The Roland–Morris disability questionnaire 12–item version.

Oswestry disability index

The original Oswestry disability index (ODI) (version 1.0)²⁰ (Table 8) includes 10 sections of questions that evaluate the activities of daily living, which can be drastically influenced by LBP. The sections have been selected from experimental questionnaires that aimed to assess several aspects of daily living. The ODI domains are the following: pain intensity, personal care, lifting, walking, sitting, standing, sleeping, sex life, social life and travelling. Each section contains six statements that are scored from 0 (minimum degree of difficulty in that activity) to 5 (maximum degree of difficulty). If more than one statement is marked in each section, the highest score should be taken. The total score is obtained by summing up the scores of all sections, giving a maximum of 50 points. The final score is expressed as a percentage with the following formula: (total score)/(5 × number of questions answered) × 100%. For example, if all 10 sections are completed the score is calculated as follows: 16 (total scored)/50 (total possible score) × 100 = 32%. If one section is missed (or not applicable) the score is calculated as follows: 16 (total scored)/45 (total possible score) × 100 = 35.5%.²¹

View this table:
[In this window](#) [In a new window](#)

Table 8
 The Oswestry disability index (version 1.0).

The authors suggest rounding the percentage to a whole number for convenience. The higher the percentage, the greater the perceived level of disability by the patient. The total score ranges from 0 to 100%, with 0 representing no disability and 100 representing maximum disability. A total score between 0 and 20% means minimal disability; between 20 and 40%, moderate disability; between 40 and 60%, severe disability; between 60 and 80% crippled; between 80 and 100%, bed bound or symptom magnifier.²⁰

The questionnaire is self–administered by the patient, it is usually completed in less than 5 min and scored in less than 1 min.

Versions of the ODI

Several versions of the ODI are available.²¹ The original version 1.0²⁰ was published without section 8 (sex life) or section 9 (social life).²² Moreover, there are two studies, in which the administration of the ODI by telephone

The Medical Research Council group produced a modified version of the ODI (version 2.0)²⁵ (Table 9), which has been proposed for general use.²⁶⁻²⁸ It has been distributed by correspondence and is available as part of a computer interview in the UK (slightly modified)^{25,29} or in the USA through MODEMS (PO Box 2354, Des Plaines, IL 60017-2354). In this version the following domains are included: pain intensity, personal care, lifting, walking, sitting, standing, sleeping, sex life (if applicable), social life and travelling. Each section contains six statements, ranging from 0 to 5, and patients should answer the questions in relation to that day ('today'). The standard scoring method can be used to obtain the final disability score.^{20,21}

View this table:
[In this window](#) [In a new window](#)

Table 9
 The Oswestry disability index (version 2.0).

A revised Oswestry Disability Questionnaire (Table 10) was published by a chiropractic study group in the UK.³⁰ This version consists of 10 sections: pain intensity, personal care, lifting, walking, sitting, standing, sleeping, social life, travelling and changing degree of pain. Also in this version each section contains six statements, ranging from 0 to 5, and the final score is calculated with standard scoring method.

View this table:
[In this window](#) [In a new window](#)

Table 10
 The revised Oswestry disability index.

A modified ODI published by Fritz and Irrgang³¹ (Table 11) is similar to the modified ODI used by Hudson-Cook *et al.*³⁰ The questionnaire consists of 10 domains: pain intensity, personal care, lifting, walking, sitting, standing, sleeping, social life, travelling and employment/homemaking. A section regarding employment and home-making ability is substituted for the section related to sex life. Each domain contains six statements, scored from 0 to 5, with higher values representing greater disability. The final score is obtained with standard scoring method.

View this table:
[In this window](#) [In a new window](#)

Table 11
 The modified Oswestry disability index.

The American Academy of Orthopaedic Surgeons (AAOS) and other spine societies have modified version 1.0 to use it like their spine outcome instrument³² (Table 12). This version includes seven sections: personal care, lifting, walking, sitting, standing, sleeping and travelling. The sections 1, 8 and 9 are omitted. Each section contains six statements, ranging from 0 to 5, and patients should answer the questions in relation to 'the past week'.¹³ The standard scoring method can be also used for this versions of the ODI, but because it has only seven sections, corrections should be made to obtain the final score.²¹

View this table:
[In this window](#) [In a new window](#)

Table 12
 AAOS/MODEMS.

Another version of the ODI has been published by the North American Spine Society (NASS).³³ This version includes a pain diagram, questions from the SF-36 health questionnaire, questions on neurological symptoms and on the LBP, and a modification of the original ODI.¹³

The Quebec back pain disability scale (QBPDS) (Table 13) is a 20-item condition-specific questionnaire to assess the degree of disability in patient with back pain.³⁴ Item selection was performed from 46 disability items by examining the test-retest reliability and responsiveness of individual items, by using techniques of factor analysis and by application of item response theory.³⁵ The QBPDS assess disability by evaluating the following daily tasks: self-care, sleeping, walking, climbing stairs, sitting, standing, lifting large or heavy objects, bending and stooping, physical activities and houseworks. Social life, sex life and pain intensity are omitted; so pain should be evaluated with other scoring system. Each one of 20 daily activities is scored with a six-point difficult scale ranging from 0 ('not difficult at all') to 5 ('unable to do'). The item scores are added up in order to obtain the disability score, which ranges between 0 and 100. The higher values represent greater disability, and sub-scores are not reported.³⁴

View this table:
[In this window](#) [In a new window](#)

Table 13
 The Quebec back pain disability scale.

The questionnaire is self-administered by the patient, it can be easily completed in about 5-10 min, and scored in less than 2 min.

The Waddell disability index

The Waddell disability index (WDI) (Table 14)³⁶ is nine-item scale which assesses disability by evaluating daily living activities commonly restricted by LBP. The items included are: lifting, sitting, standing, travelling, walking, sleeping, social life, sex life and putting on footwear. Items about work, self-care and sports are not included. Questions are not related to a specific time period and are selected from a previous questionnaire³⁷ and pilot interviews. Patients answer to questions only with positive or negative statement (yes/no). The final score is calculated by adding up positive items, and ranges from 0 to 9.³⁶

View this table:
[In this window](#) [In a new window](#)

Table 14
 The Waddell disability index.

The questionnaire is easy to administer; it can be filled out in about 5 min and scored in less than 1 min. It was validated on a chronic LBP population.

The Million visual analogue scale

The Million visual analogue scale (MVAS) (Table 15) is a 15-item questionnaire about disability and pain intensity in patients with LBP.³⁸ The 15 questions investigate the body functions (pain, sleep, stiffness and twisting), daily activities (walking, sitting, standing and work) and social life. Information about item selection process is not available. Score is given on a 100 mm visual analogue scale (VAS). For example, if patients are asked to quantify the severity of his pain (like the first question), they mark a point on a 100-mm line in which the end points are labelled as 'no pain' and 'intolerable'. In each question, it is possible to obtain an index of severity of symptoms in a patient-specific fashion measuring the distance of the marked point from the origin of the line. The final score is calculated by adding up the equally weighted scores.

View this table:
[In this window](#) [In a new window](#)

Table 15
 The Million visual analogue scale.

about 2–3 min.

The low back outcome score

The low back outcome score (LBOS)³⁹ (Table 16) is designed as a self-reported measure to assess the patients with LBP. It is a 13-item questionnaire, and it includes weighted questions about current pain, employment, domestic chores, sport activities, resting, medical treatments or consultations, drug use, sex life and daily activities (such as sleeping, walking, sitting, travelling, dressing).

View this table:
In this window In a new window

Table 16
The low back outcome score.

The pain question is answered with an 11-point VAS ranging from 'no pain' to 'maximum pain possible'. However, for scoring, the 11 answer possibilities are reduced to four categories (0–2, 3–4, 5–6, 7–10). All the other questions offer an answer for different possibilities, except the sport activities and resting questions, which provide three different answers.

The answering possibilities of each item are scored with a four-point scale, but questions are differently weighted. Three different groups of questions can be identified. Items with a nine-point scoring system (pain, employment, domestic and sport activities) in which the score can be 0, 3, 6 or 9 points. Items with a six-points scoring system (resting, treatment or consultation, analgesia and sex life) in which the score can be 0, 2, 4 or 6 points. Items with a three-points scoring system (sleeping, walking, sitting, travelling, dressing) in which the score can be 0, 1, 2 or 3 points. The final score is obtained by summing the score of each item and it ranges from 0 to 75, with lower values representing greater disability.

The questionnaire can be completed in about 5 min and scored in less than 1 min.

The low back pain rating scale

The low back pain rating scale (LBPRS)⁴⁰ (Table 17) is a rating system designed to evaluate the clinical outcome of LBP patients. This instrument includes three different components: pain, disability and physical impairment. The pain component consists of six questions divided into two groups: three questions about back pain and three questions about leg pain. Each item is scored with the VAS. Items are the following: LBP/leg pain at the time of examination (0–10 points), the worst LBP/leg pain within the last 2 weeks (0–10 points) and the average level of the back pain/leg pain during the same period (0–10 points). The final score ranges from 0 to 30 for both low back/leg pain. Therefore, the pain component in total gives 0–60 points. The disability component consists of 15 questions evaluating the patient's ability to perform daily activities, such as sleeping, ability to perform houseworks, walking, sitting, lifting, working, dressing, driving, running, getting up from a chair, climbing stairs, contact with people and expectations of future pain. Each question can be answered with three different possible answers and is scored with a three-point Likert scale. Answers are the following: 'yes' (0 points), 'can be a problem' (1 point) or 'no' (2 points). The disability component gives a total score of 0–30 points. The physical impairment component is evaluated by measuring the back muscle endurance, spinal mobility, patient mobility and use of analgesics. Muscle endurance and spinal/patient mobility are recorded with specific physical test, and each is scored on a scale ranging from 0 to 10. Use of analgesics/NSAID is scored as follows: 'no use during a week' (0 points); 'use of NSAID/non-narcotic analgesics up to 4 times a week' (2 points); 'use of NSAID/non-narcotic analgesics more than 4 times a week' (4 points); 'use of morphine/ analogues up to 4 times a week' (8 points) and 'use of morphine/analogues more than 4 times a week' (10

View this table:
[In this window](#) [In a new window](#)

Table 17
 The Low back pain rating scale.

The three different components are weighted: 60 points for pain scoring, 30 points for disability and 40 points for physical impairment. Therefore, combining them, the final LBPRS score ranges from 0 (in patient without back problems) to 130 (in disabled patient). The questionnaire can be filled out in about 15 min and scored in about 3–5 min.

The NASS lumbar spine outcome assessment instrument

The NASS lumbar spine outcome assessment instrument (NASS LSO) was first published by Daltroy *et al.*,³³ and is derived from a consensus of the NASS. It consists of 62 main question obtained from three different existing questionnaires: the SF36, a modified ODI and a modified employment assessment published by Bigos.⁴¹

The NASS data are grouped into five categories. The first group consists of demographic data (age, sex, race, education and insurance information). The second group consists of the medical history (diagnosis, past surgeries, comorbidities, etc.). The third group includes: pain, neurogenic symptoms and function. These domains are measured by a modified ODI version. The fourth group is represented by employment history, evaluated by a score system published by Bigos *et al.*⁴¹ The fifth group consists of data about outcomes of treatment, but it is included only in the follow-up module.

The scoring is complex and subscores are extractable (modified ODI, SF36, pain and disability scale, neurogenic symptoms scale, job exertion scale, expectation and satisfaction scale).⁴² The questionnaire is long and it takes 20 min to be filled out.

The clinical back pain questionnaire

The clinical back pain questionnaire (CBPQ) (Table 18), also known as the Aberdeen LBP scale, is a 19-items questionnaire, consisting of questions commonly used in the clinical assessment of patients with LBP.⁴³ It includes questions about body functions (pain, sleep, bending, loss of feeling and leg weakness) and questions about daily activities (self-care, walking, sitting, standing, sport, housework and resting). There are 6 multiple choice questions and 13 single choice questions. Answering possibilities for each question can vary between three and six items. The answer categories to each single choice question are scored in an ordinal manner (e.g. 0, 1, 2, 3 points, etc.), while multiple choice questions' responses are assigned a score of one point. The 'back pain severity score' is calculated by summing the score of the responses' to each question, and then it is converted to percentages. The final score ranges between 0 and 100 with the higher values representing greater disability.

View this table:
[In this window](#) [In a new window](#)

Table 18
 The clinical back pain questionnaire.

The questionnaire is easy to administer, can be completed within 5–10 min, and scored within 3 min.

The resumption of activities of daily living scale

The resumption of activities of daily living scale (RADL) (Table 19) scale is designed to assess broad areas often affected by back injury.⁴⁴ The scale measures the extent of resumption of a person's 'usual' activities since the time of injury. The final 12-item RADL includes the following areas: sleeping patterns, sexual activity, self-care, light and heavy household

recreational activities and paid employment. Each item is scored with a graphic scale ranging from 0 (not at all) to 100% (complete resumption). The total RADL score can vary from 0 to 100; it is obtained by summing across the items and dividing by the number of items. At least 9 items of the 12 questions have to be completed to calculate a total score for each patient.

View this table:
[In this window](#) [In a new window](#)

Table 19
 The resumption of activities of daily living scale.

The functional rating index

The functional rating index (FRI) (Table 20) is a 10-item scoring system designed to measure both patient's perception of function and pain of the spinal musculoskeletal system.⁴⁵ The instrument includes: eight items focus on daily activities (sleeping, self-care, travel, work, recreation, lifting, walking and standing) that can be affected by a spinal disease and two items focus on two different aspects of pain (intensity and frequency). Each item is scored with a five-point scale ranging from 0 (no pain or full ability to function) to 4 (worst possible pain or unable to perform a specific function at all). The index score is achieved by adding up the equally weighted scores, dividing by the maximum possible score, and multiplying by 100%. When all 10 items are answered, the formula is the following: $(\text{total score}/40) \times 100\%$. The final score ranges from 0 (representing absence of disability) to 100% (representing severe disability). Therefore, the higher the score the higher the perception of dysfunction and pain.

View this table:
[In this window](#) [In a new window](#)

Table 20
 The functional rating index.

The back pain functional scale

The back pain functional scale (BPFS) (Table 21) is a self-report measure evaluating patient's functional status in clinical and researching settings.⁴⁶ Item selection was from existing questionnaires (such as SIP,⁴⁷ OLBPD,²⁰ QBPD,³⁴ Dallas pain questionnaire (DPQ),⁴⁸ RMQ,¹⁰ MOS-36,⁴⁹ PSFS⁵⁰) and interviews with physical therapists. Items reduction was performed by examining the test-retest reliability, internal consistency, content and construct validity. The final version of the BPFS consists of 12 items, investigating work, hobbies, home activities, bending or stooping, dressing shoes or socks, lifting, sleeping, standing, walking, climbing stairs, sitting and driving. Each item is scored with a six-point scale, in which 0 means unable to perform activity, 1 extreme difficulty, 2 quite a bit of difficulty, 3 moderate difficulty, 4 a little bit of difficulty and 5 no difficulty. The total BPFS score can vary from 0, representing the lowest functional level, to 60, representing the highest functional level.

View this table:
[In this window](#) [In a new window](#)

Table 21
 The back pain functional scale.

The questionnaire takes less than 5 min to complete and about 30 s to score.

The general function score

The general function score (GFS) (Table 22) is a disease-specific instrument consisting of nine items, created to measure physical disability in patients with LBP.⁵¹ The original version consists of 17 items concerning physical activities of daily living. The final GFS includes just 9

of the 17 original items, showing high individual correlations, validity, reliability, responsiveness and feasibility. The final items are walking a flight of stairs; sitting more than 30 min; standing more than 30 min; walking more than 30 min; lifting more than 10 kg; lean over a basin; carry a bag of groceries; make the bed and dressing. Each item can be answered with three possible response alternatives: 'can perform', 'can perform with difficulty' or 'cannot perform'. These responses are respectively scored as 0, 1 and 2 points. The total score is obtained by summing each item's score, but it is represented as a percentage: 0% means no physical disability and 100% means maximal physical disability.

View this table:
[In this window](#) [In a new window](#)

Table 22
The general function score.

The questionnaire is filled out in about 2 min and scored in less than 1 min.

The patient-specific functional scale

The patient-specific functional scale (PSFS) (Table 23)⁵⁰ is a patient self-defined instrument, designed to record and measure a list of disabilities specific for each patient. The questionnaire has three separate sections: pain question set, pain limitation section and pain intensity section. The first section includes a list of activities chosen by the patient. Patients are asked to identify the five most affected activities in their daily living, because of the low back pain. A slightly different version has been also described in a population with neck dysfunction,⁵² in which the list of activities includes only three items plus the space for additional activities. To quantify the level of disability, each item is scored with an 11-point scale, ranging from 0 ('unable to perform activity') to 10 ('able to perform activity at pre-injury status level'). The second section assesses the functional limitation from pain during the previous 24 h. Pain limitation is also scored with an 11-point scale, ranging from 0 ('activities have been severely limited') to 10 ('activities have not been limited'). The third section measures pain intensity during the previous 24 h. Scoring is performed using an 11-point scale, in which the orientation is reversed, because 0 means 'no pain' and 10 means 'pain as bad as it can be'.

View this table:
[In this window](#) [In a new window](#)

Table 23
The patient-specific functional scale.

The PSFS can be administered and recorded in about 4 min. It should be administered during history taking and prior to physical examination. The clinician should read the instructions to the patient and record the activities, the corresponding score and the assessment date. At subsequent reassessments, the clinician reads the follow-up instructions and records the score of the previously identified activities and the date.⁵³

The outcome measure in lumbar spinal stenosis

The outcome measure in lumbar spinal stenosis (OMLSS) (Table 24) is a short self-administered questionnaire for the assessment of patients with lumbar spinal stenosis.⁵⁴ The questionnaire includes three sections: symptom severity, physical function and patient satisfaction. The items for each section were selected from a literature consultation and interviews with rheumatologists and orthopaedic surgeons specialized in spine surgery.

View this table:
[In this window](#) [In a new window](#)

Table 24
The outcome measure in lumbar spinal stenosis.

The symptom severity scale includes seven items: pain severity, pain frequency, pain in the back, pain in the leg, weakness, numbness and balance disturbance. Questions 1, 3, 4, 5 and 6 can be answered as follows: none, mild, moderate, severe and very severe. These responses are respectively scored with 1, 2, 3, 4 and 5 points. Also, question 2 has five possible responses scored with a five-point scale: less than once a week (1 point); at least once a week (2 points); everyday, for at least a few minutes (3 points); everyday, for most of the day (4 points) and every minute of the day (5 points). However, balance disturbance has only three answers: none (1 point), sometimes (3 points) and often (5 points). The symptom severity scale score is calculated by summing score of each answered item and dividing for the number of answered questions. The score can range from 1 to 5. If more than two items are missing, the scale score cannot be obtained.

The physical function scale consists of five questions about walking distance, ability to walk for pleasure, for shopping, for getting around the house and from bathroom to bedroom. All questions are scored with a four-point scale. Questions 2, 3, 4 and 5 can be answered as follows: yes, comfortably (1 point); yes, but sometimes with pain (2 points); yes, but always with pain (3 points) and no, could not perform (4 points). Question 1 can be answered as follows: more than 2 miles (1 point), more than 2 blocks but less than 2 miles (2 points), more than 50 feet but less than 2 blocks (3 points) and less than 50 feet (4 points). The physical function scale score is obtained by adding up score of each answered item and dividing by the number of answered questions. The score can range from 1 to 4. If more than two items are missing, the scale score cannot be calculated.

The patient satisfaction scale includes six questions about satisfaction with the overall result of the back operation, pain relief after the operation, walking ability after the operation, ability to do housework or job after the operation, strength in the thighs, legs or feet and balance or steadiness on feet. All questions are scored with a four-point scale and can be answered as follows: very satisfied (1 point), somewhat satisfied (2 points), somewhat dissatisfied (3 points) and very dissatisfied (4 points). The satisfaction scale score is obtained by summing the score of each answered item and dividing for the number of answered questions. The score can range from 1 to 4. If the number of responses exceed fours, the scale score can be calculated. The questionnaire is very easy to compile and to score.

The back illness pain and disability nine-item scale

The back illness pain and disability nine-item scale (BACKILL)⁵⁵ (Table 25) aims to detect disability and response to treatment in chronic low-back pain affected patients. Items are selected from three pre-existing validated instruments: the PAIN-FREE8, which is an 8-item version of McGill Pain Questionnaire;⁵⁶ the Functional Assessment Screening Questionnaire with five items,^{57,58} which is derived from the original 15-item FASQ;⁵⁹ the Oswestry low back pain disability questionnaire with eight items, which is a shorter version of the original OSW.²⁰ The BACKILL includes two items for pain (aching and tiring), and seven items for mobility (lifting, sitting for 30 min, standing for 30 min, travelling, getting up from a low seat, walking and personal care). Items about pain are scored with a four-point scale: none (4 points), mild (3 points), moderate (2 points) and severe (1 points). Three items about mobility (standing, sitting and getting up from a low seat) are also scored with a four-point scale: easy (4 points), a little difficulty (3 points), a lot of difficulty (2 points) and unable to do without help (1 points). Resting mobility items are scored with a six-point scale in which possible answers are specific for each question. Moreover two additional items can be included (fearful and punishing-cruel). They are scored separately from BACKILL items, with a

four-point scale. The questionnaire is self-administered and it is easy to complete and to score.

View this table:
[In this window](#) [In a new window](#)

Table 25
 The back illness pain and disability nine-item scale.

The Bournemouth questionnaire

The Bournemouth questionnaire (BQ)⁶⁰ (Table 26) is a short-form multidimensional questionnaire designed to measure the outcomes in back pain patients. The items included in the questionnaire were obtained by reviewing the literature. Seven aspects of the back pain experience were selected. These aspects were the most commonly measured, and showed significant responsiveness to clinical change. Domains are the following: pain intensity; ability to perform daily activities and social activities; anxiety status; depression status; pain interference with work activities and pain locus control. Each item is scored with an 11-point numerical rating scale from 0 to 10. A total score can be obtained by summing result of each item, although the authors recommend to express the total score of the BQ as a percentage. The questionnaire can be completed and scored quickly.

View this table:
[In this window](#) [In a new window](#)

Table 26
 The Bournemouth questionnaire.

The Dallas pain questionnaire

The DPQ⁴⁸ (Table 27) is a 16-item instrument to assess the four aspects of daily living affected by chronic back pain: day-to-day activities, such as pain and intensity, personal care, lifting, standing, sitting, walking and sleeping; work and leisure activities, such as social life, travelling and vocational; anxiety-depression status, including anxiety and mood, emotional control and depression; and social interest, such as interpersonal relationship, social support and punishing responses. Each item is scored with a VAS, divided into five, six, seven or eight small segments (it depends on the question). Scale extremities are labelled with specific words (e.g. 'no pain'/'all the time') and with percentage (0%/100%). For every specific question, the patient marks the point on the scale which represents his/her condition.

View this table:
[In this window](#) [In a new window](#)

Table 27
 The Dallas pain questionnaire.

For scoring, 0 points are assigned to the left-hand segment, 1 point to the next segment, 2 points to the next segment and so on to the last segment. Item scores are added and multiplied by a constant to obtain the percentage of pain interference with each of four daily living aspects evaluated by DPQ. The constant used for daily activities section is 3, while the constant used for work/leisure activities, anxiety/depression and social interest section is 5. The DPQ can be answered in 3–5 min and scored in less than 1 min.

The disability rating index

The disability rating index (DRI)⁶¹ (Table 28) is a 12-item questionnaire that allows to evaluate the physical function. The DRI includes the following activities: dressing; outdoor walks; climbing stairs; sitting for a longer time; standing bent over a sink; carrying a bag; making a bed; running; light work; heavy work; lifting heavy objects; participating in exercise/sports. The 12 items are divided into three categories: basic daily

life activities (questions 1–4); physical activities (questions 5–8); work-related/vigorous activities (questions 9–12).

View this table:
[In this window](#) [In a new window](#)

Table 28
The disability rating index.

Each item is scored with a 100 mm VAS. Extremities of the scale are labelled with 'without difficulty' (0 points) and 'not at all' (100 points). Patients mark a point on the line, representing their ability to perform the daily activities included in the question list. For scoring, in each item the distance in mm on the VAS between the zero points and the point marked from the patient is measured. The mean of these measurements is calculated, and the DRI score is expressed as percentage. The DRI is a very quick self-administered questionnaire, and can be scored in less than 2 min.

Jan van Breemen functional scale

Jan van Breemen functional scale (JVB)⁶² (Table 29) quantifies back pain. It consists of three different domains: pain, functional capacity and spinal mobility. The pain component includes six questions about back pain in the past week: 'in general', 'at night', 'during the first hour in the morning', 'during sitting', 'during walking' and 'during standing'. Each item is scored with an 11-point scale, ranging from 0 to 10. The functional capacity component includes eight questions about the ability to perform the following tasks during the past week: carrying, walking, standing, sitting, lifting, going outdoors, sleeping and performing household/hobby activities. Each item is also scored with a 0–10 scale. The lumbar spine mobility component is measured with the following tests: lumbar flexion index; lumbar flexion/extension index; lateral bending, fingertips to fibular head (right/left); lateral bending to the right/left, contraction; lateral bending to the right/left, distraction; active straight leg raising (right/left).

View this table:
[In this window](#) [In a new window](#)

Table 29
The Jan van Breemen functional scale.

The complete JVB (questionnaire and physical test) requires about 20 min.

The occupational role questionnaire

The occupational role questionnaire⁶³ (Table 30) is a short eight-item instrument to assess the impact of back pain in workers. It consists of two sections: productivity and satisfaction. The productivity component includes four questions about extra work, ability to work quickly, productivity/efficiency and quality of work. The satisfaction component includes four questions about opportunities to improve one's skills, job security, job satisfaction and relations with co-workers. Each of eight item is answered with four possible responses: 'a lot' (scored with 3 points); 'somewhat' (scored with 2 points); 'a little' (scored with 1 point) or 'not at all' (scored with 0 points). The final score is calculated adding up item scores and it is converted to a 0–100 scale. It is also possible to obtain two subscores, one for productivity section and one for satisfaction section. Each of them is calculated with the same format of the final ODQ score.

View this table:
[In this window](#) [In a new window](#)

Table 30
The occupational role questionnaire.

The spinal pain independence measure⁶⁴ (Table 31) is designed to assess the chronic LBP. It consists of three sections: activities related to mobility, activities performed in sitting and standing and activities performed in the room and bathroom. The mobility section includes five items: mobility for short distances, mobility for moderate distances, mobility for long distances, stair management and maximal walking speed. The activity in sitting and standing section includes three items: carrying loads, activity in the sitting position and activity in the standing position. The activity indoors section includes four items: mobility in bed, transfers, washing lower body and dressing lower body.

View this table:
[In this window](#) [In a new window](#)

Table 31
 The spinal pain independence measure.

The physical impairment scale

The physical impairment scale (PIS)⁶⁵ (Table 32) is designed to measure the physical impairment in patients with LBP. It includes seven item selected from a pool of 27 physical tests by investigating reliability, ability to discriminate patients and normal subjects, and ability to express the disability. Physical tests included in the final version of PIS are the following: total flexion, total extension, average lateral flexion, average straight leg raising (SLR), spinal tenderness, bilateral active SLR and sit-up. For each item a cut-off value is established to differentiate illness subjects from normal subjects. Each item is scored with 0 points if the test value is normal, or with 1 point if the test value is pathologic. The final score of PIS ranges from 0 to 7 and it can be expressed as percentage.

View this table:
[In this window](#) [In a new window](#)

Table 32
 The physical impairment scale.

The functional outcomes questionnaire for spinal disorders

The functional outcomes questionnaire for spinal disorders (FOQSD)⁶⁶ includes the following items: ability to perform heavy activities (such as active sports, heavy housecleaning, gardening, etc.), ability to perform light/moderate activities (such as washing, cooking, light cleaning, etc.), ability to perform activities (such as visiting friends, eating out, etc.), sitting, walking, sleeping, duration of symptoms, depression, level of pain, pain medication usage and overall satisfaction with results.

The pain response to activity and position questionnaire

The pain response to activity and position questionnaire (PRAP)⁶⁷ is a 30-item questionnaire consisting of two sections of 15 questions. One section is related to LBP and the other section is related to leg pain. In both groups of questions, for each item the patient describes his/her pain as follows: 'no pain', 'better', 'same' or 'worse'. The PRAP is a patient self-report instrument.

The back pain interference scale

The back pain interference scale⁶⁸ is an 18-item questionnaire to measure the restrictions in daily activities due to the back pain. Each question is scored with a 10-cm line scale ranging from 0 (no pain/symptoms at all) to 10 (the worst pain/symptoms). Final score vary from 0 to 180.

An analytic description for every score is presented in Table 33.

View this table:
[In this window](#) [In a new window](#)

Table 33
 Analytic description for every score.

Discussion

A wide variety of rating systems to measure functional outcomes in patients with LBP have been described in the past decades. Each of them evaluates low back performance using specific variables, including both objective and subjective criteria. Also, when the same variables are evaluated, different weight is attributed to the single domain. Interpreting these domains becomes difficult, because, even though they can be common to more than one scoring system, each stresses them in a different way. Functional status measures are usually classified as generic or disease specific.⁶⁹ Generic measures allow one to evaluate symptoms, functions or organ systems, which are not necessarily spine related; moreover they can be used in all kind of patients. Disease-specific measures assess symptoms and functional limitations related to a specific disease/condition, so in the back pain patient back-related problems are focused.⁵ Usually, physicians and researchers use both functional status measures.

Although many back pain score systems are available, the most used in clinical and research settings are: RDQ, ODI, QBPDS, WDI, MVAS, LBOS, LBPRS, NASS and CBPQ.

The RDQ is a health status measure created to assess physical disability from LBP and it is one of the most used in research or clinical settings for monitoring patients.

The RDQ is validated in English,¹⁰ French,⁷⁰ German,⁷¹ Greek,⁷² Portuguese,⁷³ Spanish,⁷⁴ Swedish,⁷⁵ Turkish,⁷⁶ Norwegian,⁷⁷ Iranian,⁷⁸ Moroccan.⁷⁹

The questionnaire is simple to complete and easily understood by patients. Patients completing the RDQ have to mark statements which describe themselves that day. The RDQ score correlate well with the data obtained from other physical function score systems, such as the QBPDS³⁴ and the ODI.²⁰ The RDQ has good construct validity, internal consistency, responsiveness and reliability.¹³ The test-retest reliability when the test-retest interval is short (24 h) is better than when the test-retest interval is long.

On the basis of the validation study conducted by Roland and Morris, the RDQ should be applied for disability assessment when there is the need to detect short-term changes in back pain or short-term changes in response to treatment. On the contrary, because of the absence of specific domains in the scale, the RDQ is inadequate when the clinicians want to assess the psychological or social problems related to the patient's LBP.¹⁰ For these reason, the RDQ should be integrated with other appropriate outcome measures when this kind of evaluation is required.

The ODI is a functional status outcome measure widely used in the clinical management of spinal disorders. It is validated in English,²⁰ Finnish,^{80,81} French,⁸² German,⁸³ Greek,⁷² Norwegian,⁷⁷ Iranian.⁷⁸ The questionnaire is quick to complete and has good construct validity, pointed out by internal consistency, responsiveness and reliability. The ODI and RDQ scores are highly correlated, with similar test-retest reliability and internal consistency.²⁵

In a recent review by Fairbank and Pynsent²¹, the authors recommend the use of ODI version 2.0 to detect meaningful changes in disability status in every day life, as when using the RDQ. However, the RDQ is recommended in patients with mild/moderate disability, whereas the ODI is recommended in patients with persistent severe disability.¹³ Moreover, unlike the RDQ, the ODI allows one to investigate the patient's social problems and sexual life.

The QBPDS is a condition specific instrument³⁴ which assesses only



functional disability and sleep, while it does not evaluate pain. The QBPDS is validated in English, Dutch,⁸⁴ French³⁵ and Iranian.⁷⁸ Because of a few validated translations to other languages, it is not as often used as the RDQ or the ODI. Internal consistency, test-retest reliability and responsiveness are satisfactory.³⁴

In the validation study by Kopec *et al.*,³⁴ the QBPDS is recommended both as outcome measurement in clinical trials and as a monitor for the patients' progresses during treatment and rehabilitation programs. Furthermore, it resulted more reliable and at least as sensitive to change when compared with the ODI and RDQ, although some authors found that the test-retest reliability and responsiveness are better for the modified ODI than the QBPDS.³¹ However, in the assessment of patients affected by LBP, the QBPDS needs to be associated with an independent pain assessment tool.⁸⁵ Moreover, it does not allow to investigate patient's social and sex life, as these items are not included in the score.

The WDI is a short nine-item score system consisting of questions about daily activities. It is validated in English,³⁶ Spanish and is also available in an unvalidated French version.⁸⁶ Internal consistency is higher in the WDI than in the RDQ, but it is lower than in the ODI.⁸⁷ Responsiveness is good: the questionnaire is sensible for clinical change 4 weeks after surgery.⁴³ Following the indications given in the validation study by Waddell and Main³⁶, in clinical settings, the WDI should be associated with other functional scoring systems to obtain a more complete assessment of disability by evaluating daily living activities commonly restricted by LBP.

The MVAS is a 15-item questionnaire evaluating disability and pain intensity in patients affected by LBP. The main advantage of MVAS scale is that it is easy to use. Despite its good reliability and internal consistency, it should be applied only when previous score are not available because it has received limited validation, and some of its questions could lead to inaccurate answers. As reported in the validation study by Million *et al.*,³⁸ the correlation between results by different observers is not always satisfactory.

The LBOS is a self-reporting measure for both assessment and outcome in patients with LBP. The LBOS should be applied when physicians need a short general assessment of current or previous back pain, medical treatments, employment, daily activities, sport activities, sex life, etc. Moreover, it is helpful in clinical settings because it is easy to administer and clearly discriminates between pain and disability.⁸⁸ Test-retest reliability is high, the internal consistency is good⁸⁹ and it correlates well with the ODI and the WDI.⁹⁰ Nevertheless, in the LBOS score pain is assessed independently and other items are scored with different scale.⁵ In this way, there is an item-weighting bias, because the total score gives different weight to questions.⁸⁴ Moreover, the LBOS is validated in English.³⁹

The LBPRS is a rating system evaluating the clinical outcome of LBP patients in clinical settings. It assesses pain, disability and physical impairment with a good internal consistency.⁴² It is available in English and validated in Danish.⁴⁰ The score is influenced by a weighting bias due to the difference in the scoring of pain (obtained with 11-point VAS scale) and scoring of all other items (obtained with a three-point Likert scale).⁸⁸ Despite its limitations, including the small number of patients recruited for the validation study, the LBPRS score is recommended in the evaluation of functional pain.⁴²

The NASS LSO is another questionnaire designed for the assessment and outcome measurements of patients with LBP. It is validated in English,³³ German⁹¹ and Italian.⁹² It represents a complete outcome assessment in which pain is a very dominant factor assessed with several measurements.

indicated clearly by using a pain locator (picture where patient has to mark the location of pain).⁸⁴ In the validation study, the authors clearly pointed out how the NASS LSO does not claim to contain the best scales at all but, given its good reliability, validity and easiness in the administration, it should be taken in consideration whenever the clinicians need to monitor patients' progress during treatment. Moreover, it allows pooling of data and it could result useful in clinical trials on LBP treatments.³³

The CBPQ, also known as the Aberdeen LBP scale, is a clinical assessment questionnaire consisting of questions about body functions and questions about daily activities. It is validated in English⁴³ and Chinese.⁹³ Internal consistency, test-retest correlation and responsiveness are acceptable.⁴³ However, the CBPQ gives different weights to the questions, pointed out by the various answering scales. In the validation study by Ruta *et al.*⁸⁸, the CBPQ scale is meant to be used in association with a general evaluation of the patient (such as the one given by SF-36) to identify health gain that enable those who treat back pain to justify their claims on scarce resources. Nevertheless, given the structural problems previously reported, the questionnaire is of limited value.

Grotle *et al.*,⁹ according to the three ICF perspectives of health (bodily, personal and social perspective), proposed a division of LBP rating systems into four groups: (i) questionnaires mainly assessing activity limitations; (ii) questionnaires mainly assessing activity limitations and few social functions; (iii) questionnaires assessing a mix of activity limitations and impairments; and (iv) questionnaires assessing items derived from all domain of functioning.

Most scores do not appear to have been constructed in a systematic fashion using recommended methodology. There is an increasing need for orthopaedic surgeons both to be familiar with and to routinely use objective measures of outcome for their procedures.⁹⁴ There is a trend towards the increased use of validated patient-based scores, but many have not been properly tested for validity, repeatability and sensitivity to change. Scores are not valid when used in a modified form and their use should be discouraged. One of the further areas of study is to compare and contrast two or more scoring scales, to ascertain whether they address the same category of low back function. In conclusion, although many scoring systems have been used to evaluate the low back function, we are still far from a single outcome evaluation system, which is reliable, valid and sensitive to clinically relevant changes, which takes into account both patients' and physicians perspective, and which is short and practical to use.

© The Author 2010. Published by Oxford University Press. All rights reserved. For permissions, please e-mail: journals.permissions@oxfordjournals.org

References

1. Fivaldo MW. Sports medicine approach to low back pain. *South Med J* 2002;95:642–646.
[Medline](#) [Web of Science](#)
2. Luo X, Pietrobon R, Sun SX, Liu CG, Hev I. Estimates and patterns of direct health care expenditures among individuals with back pain in the United States. *Spine* 2004;29:79–86.
[CrossRef](#) [Medline](#) [Web of Science](#)
3. Hart LC, Deyo RA, Cherkin DC. Physician office visits for low back pain: Frequency, clinical evaluation, and treatment patterns from a U.S. national survey. *Spine* 1995;20:11–19.
[Medline](#) [Web of Science](#)
4. Beurskens AJ, de Vet HC, Koke AJ, *et al.* Measuring the functional status of patients with low back pain. Assessment of the quality of four disease specific questionnaires. *Spine* 1995;20:1017–1028.

5. Devo RA, Andersson G, Bombardier C, et al. Outcome measures for studying patients with low back pain. *Spine* 1994;19:2032S–2036S.
[Medline](#)
6. Kopec JA, Esdaile JM. Functional disability scales for back pain. *Spine* 1995;20:1943–1949.
[Medline](#) [Web of Science](#)
7. Patrick DL, Devo RA, Atlas SJ, et al. Assessing health-related quality of life in patients with sciatica. *Spine* 1995;20:1899–1909.
[Medline](#) [Web of Science](#)
8. Koner IA. Measuring functional outcomes in persons with back pain. A review of back-specific questionnaires. *Spine* 2000;25:3110–3114.
[CrossRef](#) [Medline](#) [Web of Science](#)
9. Grötle M, Røys II, Vallestad NK. Functional status and disability questionnaires: What do they assess? A systematic review of back-specific outcome questionnaires. *Spine* 2004;30:130–140.
[Web of Science](#)
10. Roland M, Morris R. A study of the natural history of low back pain: Part 1. Development of a reliable and sensitive measure of disability in low-back pain. *Spine* 1983;8:141–144.
[Medline](#) [Web of Science](#)
11. Bergner M, Robhit RA, Kressel S, et al. The sickness impact profile: conceptual formulation and methodology for the development of a health status measure. *Int J Health Serv* 1976;6:393–415.
[Medline](#) [Web of Science](#)
12. Reurskens A, deVet HCW, Koke AIA. Responsiveness of functional status in low back pain: a comparison of different instruments. *Pain* 1996;65:71–76.
[CrossRef](#) [Medline](#) [Web of Science](#)
13. Roland M, Fairbank J. The Roland–Morris Disability Questionnaire and the Oswestry Disability Questionnaire. *Spine* 2000;25:3115–3124.
[CrossRef](#) [Medline](#) [Web of Science](#)
14. Stratford PW, Rinkley IM. Measurement properties of the RM–18: a modified version of the Roland–Morris Disability Scale. *Spine* 1997;22:2416–2421.
[CrossRef](#) [Medline](#) [Web of Science](#)
15. Dionne CE, Von Korff M, Koepsell TD, et al. A comparison of pain, functional limitations, and work status indices as outcome measures in back pain research. *Spine* 1999;24:2339–2345.
[CrossRef](#) [Medline](#) [Web of Science](#)
16. Dionne CE, Koepsell TD, Von Korff M, et al. Predicting long-term functional limitations among back pain patients in primary care settings. *J Clin Epidemiol* 1997;50:31–43.
[CrossRef](#) [Medline](#) [Web of Science](#)
17. Hinderwood MR, Barnett AC, Vickers MR. Evaluation of two time-specific back pain outcome measures. *Spine* 1999;24:1104–1112.
[CrossRef](#) [Medline](#) [Web of Science](#)
18. Walsh DA, Radcliffe IC. Pain beliefs and perceived physical disability of patients with chronic low back pain. *Pain* 2002;97:23–31.
[CrossRef](#) [Medline](#) [Web of Science](#)
19. Atlas SJ, Devo RA, van den Anker M, Singer DE, Keller RB, Patrick DL. The Maine–Seattle Back Questionnaire: a 12-item disability questionnaire for evaluating patients with lumbar sciatica or stenosis. Results of a derivation and validation cohort analysis. *Spine* 2003;28:1869–1876.
[CrossRef](#) [Medline](#) [Web of Science](#)
20. Fairbank J, Couner J, Davies J, et al. The Oswestry low back pain questionnaire. *Physiotherapy* 1980;66:271–273.
[Medline](#)

21. Fairbank JCT, Pynsent PB. The Oswestry Disability Index. *Spine* 2000;25:2940–2953.
[CrossRef](#) [Medline](#) [Web of Science](#)
22. Ronsone CR, Lawl J, Sutterlin CF. Reconstructive spinal surgery: assessment of outcome. *South Med J* 1996;89:1045–1052.
[CrossRef](#) [Medline](#) [Web of Science](#)
23. Huuli M, Sainio P, Hurri H, et al. Comparison of trunk strength measurements between two different isokinetic devices used at clinical settings. *J Spinal Disord* 1997;10:391–397.
[Medline](#) [Web of Science](#)
24. Mayo NE. Letter. *Spine* 1995;20:1535–1536.
[CrossRef](#) [Medline](#) [Web of Science](#)
25. Baker D, Pynsent P, Fairbank J. The Oswestry Disability Index revisited. In: Roland M, Jenner J, editors. *Back Pain: New Approaches to Rehabilitation and Education*. Manchester, UK: Manchester University Press; 1989. p. 174–186.
[Search Google Scholar](#)
26. Meade T, Browne W, Mallowes S, et al. Comparison of chiropractic and outpatient management of low back pain: a feasibility study. *J Epidemiol Commun Health* 1986;40:12–17.
[Abstract/FREE Full Text](#)
27. Meade TW, Dyer S, Browne W, et al. Randomized comparison of chiropractic and hospital outpatient management for low-back-pain: results from extended follow-up. *BMJ* 1995;311:349–351.
[Abstract/FREE Full Text](#)
28. Pynsent P, Fairbank J, Carr A. *Outcome Measures in Orthopaedics*. Oxford, UK: Butterworth-Heinemann; 1993.
[Search Google Scholar](#)
29. Pynsent PR, Fairbank JCT. Computer interview system for patients with back pain. *J Biomed Eng* 1989;11:25–29.
[CrossRef](#) [Medline](#) [Web of Science](#)
30. Hudson-Cook N, Tomas-Nicholson K, Breen A. A revised Oswestry disability questionnaire. In: Roland M, Jenner JB, editors. *Back Pain: New Approaches to Rehabilitation and Education*. Manchester: Manchester University Press; 1989. p. 187–204.
[Search Google Scholar](#)
31. Fritz JM, Irrgang JJ. A comparison of a modified Oswestry low back pain disability questionnaire and the Quebec back pain disability scale. *Phys Ther* 2001;81:776–788.
[Abstract/FREE Full Text](#)
32. Fairbank J. Use of Oswestry disability index (ODI). *Spine* 1995;20:1535–1537.
[CrossRef](#) [Medline](#) [Web of Science](#)
33. Daltroy LH, Cate Riril WL, Katz IN, Fossel AH, Liang MH. The North American spine society lumbar spine outcome assessment instrument: reliability and validity tests. *Spine* 1996;21:741–749.
[CrossRef](#) [Medline](#) [Web of Science](#)
34. Konec IA, Fedaille IM, Abrahamowicz M, et al. The Quebec back pain disability scale: measurement properties. *Spine* 1995;20:341–352.
[Medline](#) [Web of Science](#)
35. Konec IA, Fedaille IM, Abrahamowicz M, et al. The Quebec back pain disability scale: conceptualization and development. *J Clin Epidemiol* 1996;49:151–161.
[CrossRef](#) [Medline](#) [Web of Science](#)
36. Waddell G, Main CJ. Assessment of severity in low-back disorders. *Spine* 1984;9:204–208.
[CrossRef](#) [Medline](#) [Web of Science](#)
37. Wing PC, Wilfling FI, Kokan PI. Psychological demographic and orthopaedic

- 1973;90:153–160.
[Medline](#)
38. Million R, Hall W, Nilzen KH, Baker RD, Jansson MI. Assessment of the progress of the back-pain patient. 1981 Volvo Award in Clinical Science. *Spine* 1981;7:204–212.
[Search Google Scholar](#)
39. Greenough CC, Fraser RD. Assessment of outcome in patients with low-back pain. *Spine* 1992;17:36–41.
[Medline](#) [Web of Science](#)
40. Manniche C, Asmussen K, Lauritsen R, et al. Low back pain rating scale: validation of a tool for assessment of low back pain. *Pain* 1994;57:317–326.
[CrossRef](#) [Medline](#) [Web of Science](#)
41. Rinos SI, Battie MC, Snøden DM, et al. A prospective study of work perceptions and psychosocial factors affecting the report of back injury. *Spine* 1991;16:1–6.
[CrossRef](#) [Medline](#) [Web of Science](#)
42. Müller H, Duetz MS, Roeder C, Greenough CC. Condition-specific outcome measures for low back pain. Part I: Validation. *Eur Spine J* 2004;13:301–313.
[Medline](#) [Web of Science](#)
43. Ruta DA, Carratt AM, Wardlaw D, et al. Developing a valid and reliable measure of health outcome for patients with low back pain. *Spine* 1994;19:1887–1896.
[Medline](#) [Web of Science](#)
44. Williams RM, Myers AM. New approach to measuring recovery in injured workers with acute low back pain: resumption of activities of daily living scale. *Phys Ther* 1998;78:613–623.
[Abstract/FREE Full Text](#)
45. Feise RJ, Menke IM. Functional rating index: A new valid and reliable instrument to measure the magnitude of clinical change in spinal conditions. *Spine* 2001;26:78–87.
[CrossRef](#) [Medline](#) [Web of Science](#)
46. Stratford PW, Rinklev IM, Riddle DI. Development and initial validation of the back pain functional scale. *Spine* 2000;25:2095–2102.
[CrossRef](#) [Medline](#) [Web of Science](#)
47. Bergner M, Robbitt RA, Carter WR, Gilson BS. The sickness impact profile: development and final revision of a health status measure. *Med Care* 1981;19:787–805.
[Medline](#) [Web of Science](#)
48. Lawlis GF, Cuevas R, Selby D, McCoy CE. The development of the Dallas Pain Questionnaire: an assessment of the impact of spinal pain on behavior. *Spine* 1989;14:511–516.
[CrossRef](#) [Medline](#) [Web of Science](#)
49. Ware JE, Jr, Sherbourne CD. The MOS 36-item short-form health survey (SF-36) I. Conceptual framework and items selection. *Med Care* 1992;30:473–483.
[Medline](#) [Web of Science](#)
50. Stratford PW, Gill C, Westaway M, Rinklev I. Assessing disability and change on individual patients: a report of a patient specific measure. *Physiother Can* 1995;47:258–263.
[CrossRef](#)
51. Hägg G, Fritzell P, Romberg K, Nordwall A. The General Function Score: a useful tool for measurement of physical disability. Validity and reliability. *Eur Spine J* 2001;10:203–210.
[CrossRef](#) [Medline](#) [Web of Science](#)
52. Westaway MD, Stratford PW, Rinklev IM. The patient-specific functional scale: validation of its use in persons with neck dysfunction. *J Orthop Sports Phys Ther* 1998;27:331–338.
[CrossRef](#) [Medline](#) [Web of Science](#)
53. Chatman AB, Hyams SP, Neel IM, et al. The Patient-Specific Functional Scale:

- 1997;77:820–829.
[Search Google Scholar](#)
54. Stucki G, Daltroy L, Liang MH, Linton SJ, Fossel AH, Katz IN. Measurement properties of a self-administered outcome measure in lumbar spinal stenosis. *Spine* 1996;21:796–803.
[CrossRef](#) [Medline](#) [Web of Science](#)
55. Tesin I, Granger CV, Fiedler RC. A unidimensional pain/disability measure for low-back pain syndromes. *Pain* 1997;69:269–278.
[CrossRef](#) [Medline](#) [Web of Science](#)
56. Melzack R. The short-form McGill Pain Questionnaire. *Pain* 1987;30:191–197.
[CrossRef](#) [Medline](#) [Web of Science](#)
57. Granger CV, Wright RD. Looking ahead to the use of functional assessment in ambulatory physiatric practice. *Phys Med Rehabil Clin North Am* 1993;4:595–605.
[Search Google Scholar](#)
58. Granger CV, Ottenbacher KI, Baker IC, Sehgal A. Reliability of a brief outpatient functional outcome assessment measure. *Am J Phys Med Rehabil* 1995;74:469–475.
[CrossRef](#) [Medline](#) [Web of Science](#)
59. Seltzer GR, Granger CV, Wineberg DF. Functional assessment: bridge between family and rehabilitation medicine within an ambulatory practice. *Arch Phys Med Rehabil* 1982;63:453–457.
[Medline](#) [Web of Science](#)
60. Bolton JE, Breen AC. The Bournemouth Questionnaire: a short-form comprehensive outcome measure. Psychometric properties in back pain patients. *J Manipulative Physiol Ther* 1999;22:503–510.
[CrossRef](#) [Medline](#) [Web of Science](#)
61. Salén RA, Spangfort EV, Nvåren AI, Nordemar R. The Disability Rating Index: an instrument for the assessment of disability in clinical settings. *J Clin Epidemiol* 1994;47:1423–1434.
[CrossRef](#) [Medline](#) [Web of Science](#)
62. Iankhorst GI, Van de Stadt RJ, Vonelaar TW, et al. Objectivity and repeatability of measurements in low back pain. *Scand J Rehab Med* 1982;14:21–26.
[CrossRef](#) [Medline](#) [Web of Science](#)
63. Konec JA, Fedala IM. Occupational role performance in persons with back pain. *Dis Rehabil* 1998;20:373–379.
[CrossRef](#)
64. Itzkovich M, Catz A, Tamir A, et al. Spinal pain independence measure—a new scale for assessment of primary ADL dysfunction related to LBP. *Dis Rehabil* 2001;23:186–191.
[CrossRef](#)
65. Waddell G, Somerville D, Henderson I, Newton M. Objective clinical evaluation of physical impairment in chronic low back pain. *Spine* 1992;17:617–628.
[Medline](#) [Web of Science](#)
66. Nork SF, Hu SS, Workman KI, et al. Patient outcomes after decompression and instrumented posterior spinal fusion for degenerative spondylolisthesis. *Spine* 1999;24:561–569.
[CrossRef](#) [Medline](#) [Web of Science](#)
67. Roach KE, Brown MD, Albin RD, et al. The sensitivity and specificity of pain response to activity and position in categorizing patients with low back pain. *Phys Ther* 1997;77:730–738.
[Abstract/FREE Full Text](#)
68. Rytokoski H, Puukka P, Talo S. Invalidities in relation to other disease consequences in the functional assessment of patients with chronic low back pain. *Int J Rehabil Res* 1997;20:225–244.
[Search Google Scholar](#)

69. Patrick DL, Devn RA. Generic and disease specific measures in assessing health status and quality of life. *Med Care* 1989;27(Suppl.):S217–S232.
[Medline](#) [Web of Science](#)
70. Coste J, Le Parc JM, Berge F, Delcroix J, Paolaggi IR. French validation of a disability rating scale for the evaluation of low back pain (EIFEL questionnaire). 1993;60:335–341.
[Search Google Scholar](#)
71. Wiasinger GE, Nuhr M, Quittan M, Ebenbichler G, Wolf G, Fialka Moser V. Cross-cultural adaptation of the Roland–Morris questionnaire for German-speaking patients with low back pain. *Spine* 1999;24:1099–1103.
[CrossRef](#) [Medline](#) [Web of Science](#)
72. Roscaino PI, Sankas G, Stilianesi F, Prouskas K, Panagakis SA. Greek versions of the Oswestry and Roland–Morris disability questionnaires. *Clin Orthop* 2003;40–53.
[Search Google Scholar](#)
73. Nuschau I, Natour I, Ferraz MB, Goldenberg I. Translation, adaptation and validation of the Roland–Morris questionnaire. *Brasil: Bol Lett Med Biol Bras – Revista Brasileira de pesquisas médicas e biológicas/Sociedade Brasileira de Biofísica...* [et al] 2001;34:203–210.
[Search Google Scholar](#)
74. Kovacs EM, Iñohera I, Gil Del Real MT, et al. Validation of the Spanish version of the Roland–Morris questionnaire. *Spine* 2002;27:538–542.
[CrossRef](#) [Medline](#) [Web of Science](#)
75. Johansson E, Lindberg P. Subacute and chronic low back pain. Reliability and validity of a Swedish version of the Roland and Morris Disability Questionnaire. *Scand J Rehabil Med* 1998;30:139–143.
[CrossRef](#) [Medline](#) [Web of Science](#)
76. Kurukdeveci AA, Tennant A, Elhan AH, Nivazoglu H. Validation of the Turkish version of the Roland–Morris Disability Questionnaire for use in low back pain. *Spine* 2001;26:2738–2743.
[CrossRef](#) [Medline](#) [Web of Science](#)
77. Grøtli M, Brøy H, Vallestad NK. Cross-cultural adaptation of the Norwegian versions of the Roland–Morris Disability Questionnaire and the Oswestry Disability Index. *J Rehabil Med* 2003;35:241–247.
[CrossRef](#) [Medline](#) [Web of Science](#)
78. Mousavi SI, Parniannour M, Mehdian H, Montazeri A, Mohini R. The Oswestry Disability Index, the Roland–Morris disability questionnaire, and the Quebec back pain disability scale: translation and validation studies of the Iranian versions. *Spine* 2006;31:E454–E459.
[CrossRef](#) [Medline](#) [Web of Science](#)
79. Maaroufi H, Benhouazza K, Faik A, et al. Translation, adaptation, and validation of the Moroccan version of the Roland–Morris Disability Questionnaire. *Spine* 2007;32:1461–1465.
[CrossRef](#) [Medline](#) [Web of Science](#)
80. Grönlund M, Hunli M, Wannerstrand P, et al. Intercorrelation and test–retest reliability of the pain disability index (PDI) and the Oswestry disability questionnaire (ODQ) and their correlation with pain intensity in low back pain patients. *Clin J Pain* 1993;9:189–195.
[Medline](#) [Web of Science](#)
81. Grönlund M, Jarvinen E, Hurri H, Hunli M, Karaharju EO. Relationship of the Pain Disability Index (PDI) and the Oswestry Disability Questionnaire (ODQ) with three dynamic physical tests in a group of patients with chronic low-back and leg pain. *Clin J Pain* 1994;10:197–203.
[Medline](#) [Web of Science](#)
82. Dronov B, Marty M. Quality-of-life indexes for assessment of low back pain. *Rev Rhum* 1994;61:S44–S48.
[Web of Science](#)
83. Basler H, Jakle C, Kroner–Herwin B. Incorporation of cognitive behavioral

- randomized study in German pain treatment centers. *Patient Educ Counsel* 1997;31:113–124.
[CrossRef](#) [Medline](#) [Web of Science](#)
84. Schönnink JF, van Tulder MW, Koes RW, Reurkens SA, de Rie RA. Reliability and validity of the Dutch adaptation of the Quebec Back Pain Disability Scale. *Phys Ther* 1996;76:268–275.
[Abstract/FREE Full Text](#)
85. Müller H, Röder C, Greenough CG. Back related outcome assessment instruments. *Eur Spine J* 2006;15:S25–S31.
[CrossRef](#) [Medline](#) [Web of Science](#)
86. Guillemin F, Constant F, Collin JF, Boulangé M. Short and longterm effect of spa therapy in chronic low back pain. *Br J Rheumatol* 1994;33:148–151.
[Abstract/FREE Full Text](#)
87. Davidson M, Keating JI. A comparison of five low back disability questionnaires: reliability and responsiveness. *Phys Ther* 2002;82:8–24.
[Abstract/FREE Full Text](#)
88. Müller H, Röder C, Dubs I, Dietz MS, Greenough CG. Condition-specific outcome measures for low back pain. Part II: Scale construction. *Eur Spine J* 2004;13:314–324.
[CrossRef](#) [Medline](#) [Web of Science](#)
89. Holt AE, Shaw NL, Shetty A, Greenough CG. The reliability of the low back outcome score for back pain. *Spine* 2002;27:206–210.
[CrossRef](#) [Medline](#) [Web of Science](#)
90. Taylor SJ, Taylor AE, Foy MA, et al. Responsiveness of common outcome measures for patients with low back pain. *Spine* 1999;24:1805–1812.
[CrossRef](#) [Medline](#) [Web of Science](#)
91. Pace R, Sangha O, Peters A, Wildner M. Validation of the North American Spine Society instrument for assessment of health status in patients with chronic backache. *Z Orthop Ihre Grenzgeb* 1999;137:437–441.
[CrossRef](#) [Medline](#) [Web of Science](#)
92. Padua R, Padua I, Ceccarelli F, et al. Cross-cultural adaptation of the lumbar North American Spine Society questionnaire for Italian-speaking patients with lumbar spinal disease. *Spine* 2001;26:E344–E347.
[CrossRef](#) [Medline](#)
93. Leung AS, Lam TH, Hedley AJ, Twomey JT. Use of a subjective health measure on Chinese low back pain patients in Hong Kong. *Spine* 1999;24:961–966.
[CrossRef](#) [Medline](#) [Web of Science](#)
94. Longo UG, Franceschi F, Lonnini M, Maffulli N, Denaro V. Rating systems for evaluation of the elbow. *Br Med Bull* 2008;87:131–161.
[Abstract/FREE Full Text](#)

Related articles

Editor's Choice:

Norman Vetter

Editor's Choice

Br Med Bull (2010) 94 (1): 1–5 doi:10.1093/bmb/ldq013

[Extract](#) [Free](#) [Full Text \(HTML\)](#) [Full Text \(PDF\)](#)

Articles citing this article

Online Systematic Review of Single versus Double Bundle Anterior Cruciate Ligament Reconstruction

Copyright © 2013 Oxford University Press

Br Med Bull (2011) 97 (1): 47-80

[Abstract](#) [Full Text \(HTML\)](#) [Full Text \(PDF\)](#)

[Site Map](#) [Privacy Policy](#) [Cookie Policy](#) [Legal Notices](#) [Frequently Asked Questions](#)

Other Oxford University Press sites:

