






BMJ Open Clinical outcome measures and their evidence base in degenerative cervical myelopathy: a systematic review to inform a core measurement set (AO Spine RECODE-DCM)

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ABSTRACT

Objectives To evaluate the measurement properties of outcome measures currently used in the assessment of degenerative cervical myelopathy (DCM) for clinical research.

Design Systematic review

Data sources MEDLINE and EMBASE were searched through 4 August 2020.

Eligibility criteria Primary clinical research published in English and whose primary purpose was to evaluate the measurement properties or clinically important differences of instruments used in DCM.

Data extraction and synthesis Psychometric properties and clinically important differences were both extracted from each study, assessed for risk of bias and presented in accordance with the Consensus-based Standards for the selection of health Measurement Instruments criteria.

Results Twenty-nine outcome instruments were identified from 52 studies published between 1999 and 2020. They measured neuromuscular function (16 instruments), life impact (five instruments), pain (five instruments) and radiological scoring (five instruments). No instrument had evaluations for all 10 measurement properties and <50% had assessments for all three domains (ie, reliability, validity and responsiveness). There was a paucity of high-quality evidence. Notably, there were no studies that reported on structural validity and no high-quality evidence that discussed content validity. In this context, we identified nine instruments that are interpretable by clinicians: the arm and neck pain scores; the 12-item and 36-item short form health surveys; the Japanese Orthopaedic Association (JOA) score, modified JOA and JOA Cervical Myelopathy Evaluation Questionnaire; the neck disability index; and the visual analogue scale for pain. These include six scores with barriers to application and one score with insufficient criterion and construct validity.

Strengths and limitations of this study

- Consensus-based reporting guidelines were used to evaluate the properties and clinically important differences of degenerative cervical myelopathy measurement instruments.
- Only instruments that are currently in use were evaluated in this study.
- Interpretability was used as an important characteristic to make recommendations, a posteriori, due to the absence of category A Consensus-based Standards for the selection of health Measurement Instruments (COSMIN) recommendations.
- Interpretability and feasibility were evaluated using bespoke criteria adapted, a priori, from the COSMIN methodology.

Conclusions This review aggregates studies evaluating outcome measures used to assess patients with DCM. Overall, there is a need for a set of agreed tools to measure outcomes in DCM. These findings will be used to inform the development of a core measurement set as part of AO Spine RECODE-DCM.

INTRODUCTION

The most common adult spinal cord disease, degenerative cervical myelopathy (DCM), is both measured and reported inconsistently across clinical research.^{1–4} DCM is a progressive spinal cord disease caused by degenerative changes in the cervical spine that lead to stress and injury to the cervical spinal cord. It usually initially presents as a loss of digital dexterity, subtle gait disturbances and mild

pain which, if left untreated, can potentially lead to tetraplegia and wheelchair dependence.⁵

In 2019, AO Spine launched the Research Objectives and Common Data Elements for Degenerative Cervical Myelopathy (AO Spine RECODE-DCM; www.aospine.org/recode) initiative with the aim of creating a 'research toolkit' to help accelerate knowledge discovery and improve outcomes in DCM.^{3 6} The initiative identified the need to improve consistency in measurement and reporting across DCM research to enable studies to be compared and/or aggregated, and to ensure the most meaningful aspects of the disease are captured.^{7 8} This process started by creating a list of essential outcomes (ie, core outcome set) and baseline characteristics (ie, core data elements). To truly enable consistent reporting, however, these datasets should be partnered with a core measurement set (CMS): a set of agreed tools that are used to measure the outcomes and data elements of DCM.^{9–17}

Several approaches have been employed to form a CMS, ranging from the development of novel measurement instruments to adopting the use of existing ones.^{18–20} For AO Spine RECODE-DCM, it was decided to recommend existing instruments and, preferably, those already used in DCM. This was to allow a more rapid introduction of the CMS, cognisant that many new tools are in development and the CMS can be updated in the future.

Consequently, we sought to examine the tools used in DCM research and assess their quality²¹ using objective criteria. In recognition of variable quality among reported outcome measures, the Consensus-based Standards for the selection of health Measurement Instruments (COSMIN) initiative has developed clinimetric tools to assess instrument quality.²² We searched the literature for studies evaluating one or more psychometric properties defined by the COSMIN guidelines, as well as studies that defined clinically important differences such as the minimally clinical important difference (MCID) and substantial clinical benefits (SCBs). Data were rated, aggregated and assessed for methodology bias using the COSMIN manual for systematic reviews of patient-reported outcome measures (PROMs).^{23–25} This work builds on the protocol for the AO Spine RECODE-DCM initiative^{3 6 26} and complements two earlier reviews of outcome measures in DCM.^{2 21}

METHODS

Search

A search string was developed to identify original research assessing the psychometric properties of instruments currently used in the clinical research of DCM.²⁷ This comprised synonyms of 'psychometric' and 'DCM' (online supplemental table 1). The search was developed with oversight of a medical librarian (IK) and informed by previously developed search filters for DCM.^{27–29} The search was applied to MEDLINE and EMBASE, from inception until 4 August 2020, using OVID (Wolters

Table 1 Inclusion and exclusion criteria

Inclusion	Exclusion
Publication type	
<ul style="list-style-type: none"> ▶ Article written in English ▶ Primary clinical research articles 	<ul style="list-style-type: none"> ▶ Article not written in English ▶ Conference abstracts or posters ▶ Editorials, commentaries, opinion papers or letters ▶ Book chapters or theses
Study type	
<ul style="list-style-type: none"> ▶ Study includes primary clinical data 	<ul style="list-style-type: none"> ▶ Study uses only secondary data ▶ Case reports ▶ Narrative reviews ▶ Systematic reviews ▶ Meta-analyses
Populations	
<ul style="list-style-type: none"> ▶ Human studies 	<ul style="list-style-type: none"> ▶ Non-human studies
Indications	
<ul style="list-style-type: none"> ▶ Exclusively DCM (CSM, OPLL, cervical stenosis, spondylosis, spinal cord compression, cervical myelopathy) 	<ul style="list-style-type: none"> ▶ Populations with DCM and at least one other condition (eg, radiculopathy)
Comparator	
<ul style="list-style-type: none"> ▶ At least one assessment tool^{2 21 30} 	
Outcomes	
<ul style="list-style-type: none"> ▶ At least one psychometric property ▶ At least one MCID or SCB 	
<p>CSM, Cervical spondylotic myelopathy; DCM, degenerative cervical myelopathy; MCID, minimally clinical important difference; OPLL, Ossification of the posterior longitudinal ligament; SCB, substantial clinical benefits.</p>	

Kluwer, Netherlands). The search also focused on DCM tools identified in previous scoping reviews.^{2 21 30}

Study selection

All titles and abstracts were screened independently against a set of predefined eligibility criteria by four reviewers (AYT, AB, ED and FB). A full list of inclusion and exclusion criteria of studies are stated in [table 1](#).

Potentially eligible studies were selected for full-text analysis. In the event of multiple publications analysing the same cohort for the same purpose, the most recent paper was used for evaluation. At each stage, two reviewers independently (AYT, AB, ED, FB) reviewed all the screened studies for inclusion to ensure reliability of study selection (online supplemental table 2). Disagreements were resolved by consensus or appeal to a third senior reviewer (BMD).

Table 2 Definitions of domains, measurement properties and aspects of measurement properties, adapted from the COSMIN guidelines^{23–25 48} and studies of clinically important differences^{49 50}

Domain	Measurement property	Aspect	Definition
Reliability			The degree to which the measurement is free from measurement error. The extent to which scores for patients who have not changed are the same for repeated measurement under several conditions; over time; by different persons on the same occasion; or by the same persons on different occasions.
	Internal consistency		The degree of inter-relatedness among the items included in a measurement instrument.
	Reliability		The proportion of the total variance in the measurements which is due to ‘true’* differences between patients.
	Measurement error		The systematic and random error of a patient’s score that is not attributed to true changes in the construct to be measured.
Validity			The degree to which a measurement instrument measures the construct(s) it purports to measure.
	Content validity		The degree to which the content of a measurement tool is an adequate reflection of all facets of a given construct.
	Construct validity		The degree to which the scores of a measurement instrument are consistent with hypotheses (for instance, with regard to internal relationships, relationships to scores of other instruments or differences between relevant groups) based on the assumption that the instrument validly measures the construct to be measured.
		Structural validity	The degree to which the scores of a measurement instrument are an adequate reflection of the dimensionality of the construct to be measured.
		Hypotheses testing	Idem construct validity.
		Cross-cultural validity	The degree to which the performance of the items on a translated or culturally adapted measurement instrument are an adequate reflection of the performance of the items of the original version of the instrument.
	Criterion validity		The degree to which the scores of a measurement instrument are an adequate reflection of a ‘gold standard’.
Responsiveness			The ability of a measurement instrument to detect change over time in the construct to be measured.
	Responsiveness		Idem responsiveness.
Interpretability†			Interpretability is the degree to which one can assign qualitative meaning—that is, clinical or commonly understood connotations—to a PROM’s quantitative scores or change in scores.
	Clinically important differences		
		Minimal clinically important difference	The smallest measured change score that patients perceive to be important, also known as the MCID or MID
		Substantial clinical benefit	The change in outcome associated with patient perception of a large meaningful improvement.

*The word ‘true’ must be seen in the context of the classical test theory, which states that any observation is composed of two components—a true score and error associated with the observation. ‘True’ is the average score that would be obtained if the scale was applied infinite number of times. It refers only to the consistency of the score, and not to its accuracy.⁵¹

†Interpretability is not considered a measurement property, but an important characteristic of a measurement instrument.

COSMIN, Consensus-based Standards for the selection of health Measurement Instruments; MCID, minimally clinical important difference; MID, minimally important difference; PROMs, patient-reported outcome measures.

Quality assessment

The quality of included studies was assessed using the COSMIN risk of bias checklist.^{23–25} Briefly, the COSMIN risk of bias tool assesses 10 measurement properties, including nine psychometric properties (ie, content validity, structural validity, internal consistency, cross-cultural validity/measurement invariance, reliability, measurement error, criterion validity, hypotheses testing for construct validity and responsiveness) and clinically important differences. A list of definitions is presented in table 2. Interpretability and feasibility were also evaluated using criteria adapted a priori from the COSMIN

methodology (online supplemental tables 3 and 4), respectively). Namely, interpretability was evaluated for each measurement instrument through the availability of anchor-based MCIDs,^{23–25} while feasibility was assessed with respect to the ease of application of the instrument.

The methodological quality of each study was scored as ‘very good’, ‘adequate’, ‘doubtful’, ‘inadequate’ or ‘not applicable’. Overall ratings were then made for each property using the modified Grading of Recommendations Assessment, Development, and Evaluation approach from the COSMIN risk of bias checklist.^{23–25} For each study, one review author (AYT) assessed the quality,

feasibility and interpretability from included studies and a second (BD) checked the assessments. Disagreements were resolved by consensus.

Data extraction

A proforma adapted from COSMIN was employed by one reviewer (AYT) to extract the following: study details, sample size, patient demographics, measurement properties and qualitative and/or quantitative results for each property. This was checked by a second reviewer (BD) and any disagreements were resolved by consensus. Examples of qualitative and quantitative results included observations (eg, narrative syntheses) and statistics (eg, correlation coefficients). These result types are specific for each measurement property and are listed in the COSMIN guidelines.^{23–25}

Data analysis

Each result was rated as ‘sufficient’, ‘indeterminate’ or ‘insufficient’. All results were qualitatively summarised and given an overall rating as ‘sufficient’, ‘indeterminate’, ‘inconsistent’ or ‘insufficient’. The definitions of these ratings are available in the COSMIN guidelines.^{23–25} Measurement instruments were categorised into three recommendation groups:

1. Instruments with evidence of sufficient content validity and at least low-quality evidence of sufficient internal consistency.
2. Instruments categorised not in 1 or 3.
3. Instruments with high-quality evidence of an insufficient measurement property.^{23–25}

Recommendations for each instrument were presented in tandem with interpretability and feasibility assessments and reported as a narrative synthesis.³¹ We used the Preferred Reporting Items for Systematic Reviews and Meta-Analyses checklist when writing our report.³²

Patient and public involvement

This project forms part of a larger, international multi-stakeholder co-production initiative called AO Spine RECODE-DCM, which aims to develop a framework to accelerate knowledge discovery that can improve outcomes in DCM. Patients and the public were therefore involved in its overall design, conduct, management, and dissemination, and are recognised among the authors of this article. For further information, please refer to www.aospine.org/recode.

RESULTS

Literature search

The primary literature search identified a total of 3239 unduplicated studies (MEDLINE: 2389, EMBASE: 1550). Abstract and full-text screening excluded 3187 studies. Therefore, this review included a total of 52 studies (figure 1 and online supplemental table 2).

Study properties

The 52 included studies assessed a total of 7395 patients worldwide (female: 3217, male: 4178) with 29 instruments

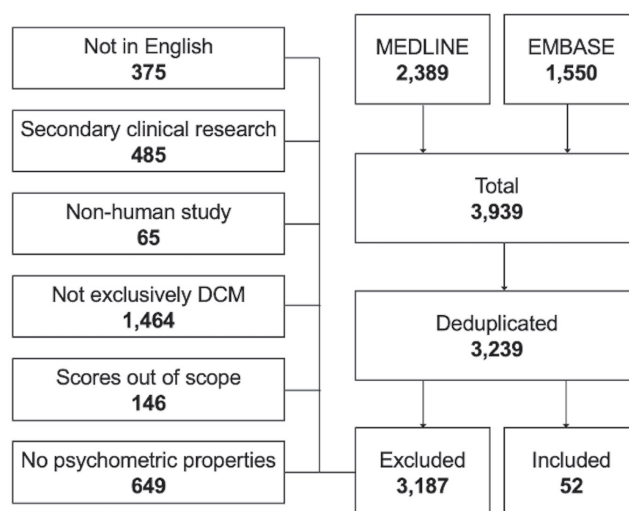


Figure 1 Preferred Reporting Items for Systematic Reviews and Meta-Analyses flow chart. A systematic review of Medline and EMBASE was conducted through 4 August 2020 to identify original research on the measurement properties of instruments currently used in degenerative cervical myelopathy research.

(table 3). These were classified into four domains based on the DCM core outcome set³³: neuromuscular function, life impact, pain, and radiological scoring.

Measurement properties

The measurement properties of the 29 instruments were evaluated using the COSMIN methodology for systematic reviews.^{23–25} A summary of findings is presented in table 4¹: the overall feasibility rating,² the overall interpretability rating and³ the overall recommendation category based on existing evidence. Included studies reported on at least one of the 10 COSMIN properties for all instruments. No instrument had evidence for all 10 properties and <50% (13/29) of instruments had evidence for at least one property per measurement domain (figure 2).

Content validity

Only three measurement instruments were evaluated for content validity: the JOA Cervical Myelopathy Evaluation Questionnaire (JOACMEQ), the modified JOA (mJOA) score and the Berg Balance Scale (BBS) (online supplemental table 5). The overall ratings for content validity, however, were indeterminate due to the uncertainty of the methods used to assess comprehensibility, and the very low quality of the evidence.

Structural validity

No instruments were assessed for structural validity.

Internal consistency

Ten measurement instruments were evaluated for internal consistency, including the JOACMEQ, JOA, mJOA, 12-Item Short Form Health Survey (SF-12) and SF-36 (online supplemental table 6). Since structural validity is required for the interpretation of internal consistency,

Table 3 Study properties

Property	Number	%
Total studies, included	52	100
Prospective	31	60
Retrospective	21	40
Total patient sample	7395	100
Male	4178	56
Female	3217	44
Measurement instruments by domain*	29	100
Neuromuscular function	16	55
Life impact	5	17
Pain	5	17
Radiological scoring	5	17
Publication year		
Maximum year of publication	2020	–
Median year of publication	2014	–
Mean year of publication	2012	–
Minimum year of publication	1999	–
Countries, by number of patients	7395	100
Japan	2014	27
USA	1802	24
Canada	1361	18
South Korea	726	10
Global/multicentre	601	8
China	255	3
India	121	2
Iran	87	1
Brazil	85	1
Italy	75	1
Hong Kong	72	1
Thailand	70	1
Taiwan	45	1
UK	41	1
France	40	1

*Instrument counts per domain do not add up to the total due to the one-to-many relationship between certain instruments and domains (eg, JOACMEQ is used both for life impact and neuromuscular function; see [table 4](#)).

JOACMEQ, Japanese Orthopaedic Association Cervical Myelopathy Evaluation Questionnaire.

the overall ratings for internal consistency were indeterminate, given the aforementioned absence of studies on structural validity.

Cross-Cultural validity

Only three measurement instruments were evaluated for cross-cultural validity: JOACMEQ, JOA and mJOA (online supplemental table 7). The overall ratings were indeterminate due to the absence of multiple group factors

analyses and differential item functioning analyses. The quality of evidence was also very low due to the uncertainty of the approaches used to analyse the data.

Reliability

Seventeen measurement instruments were evaluated for reliability, including JOACMEQ, JOA and mJOA (online supplemental table 8). The reported measures of reliability were test–retest reliability, intraobserver reliability and interobserver reliability. No instrument attained high-quality evidence for sufficient or insufficient reliability due to (1) imprecision (sample sizes <100), (2) serious inconsistency and/or (c) serious risk of bias.

Measurement error

Nine instruments were evaluated for measurement error, including JOACMEQ, JOA, mJOA, NDI, SF-36 and Visual Analogue Scale (VAS) for pain (online supplemental table 9)). The measures of error reported were minimal detectable change and distribution-based MCID.^{23–25 34} The mJOA was the only score to attain high-quality evidence for sufficiency (distribution-based MCID range: 1.2–1.4, total sample size: 868). Due to the inconsistency of results, the quality of the evidence of most other instruments could not be rated.

Criterion validity

Twelve measurement instruments were evaluated for criterion validity, including the JOACMEQ, JOA, mJOA, NDI and SF-36 (online supplemental table 10). Both the mJOA and the patient-derived version of the mJOA (P-mJOA) attained high-quality evidence for sufficient criterion validity as whole scales. However, three of four items of the mJOA, along with the 10 s step test and foot tapping test, attained high-quality evidence for insufficient criterion validity (ie, these subdomains lack criterion validity for their use as separate measures). The quality of the evidence of most of the remaining instruments was not high due to (1) imprecision (ie, sample sizes <100) or (b) important methodological flaws in the design or statistical methods.

Construct validity

Sixteen measurement instruments were evaluated for construct validity, including JOACMEQ, JOA, mJOA, NDI, arm and neck pain scores and SF-12 (online supplemental table 11). From these, 8 of 16 attained high-quality evidence for sufficient construct validity; these included the NDI, arm and neck pain scores and SF-12. Two instruments achieved high-quality evidence for insufficient construct validity. Notably, the mJOA had both high-quality sufficiency and insufficiency depending on the comparator tool (eg, sufficiency with respect to the NDI and SF-36 and insufficiency with respect to the 30 m walking test (30MWT) and EuroQol-5 Dimension (EQ-5D)). While the designs and statistical methods applied were adequate for the research questions posed, the quality of the evidence of most of the remaining tools ranged from ‘low’ to ‘moderate’ due to imprecision (ie,

**Table 4** Summary of findings

Domain	Instrument	Feasibility	Interpretability	Recommendation category	Recommendation justification
Life impact					
	EQ-5D	+	+	C	High-quality evidence for insufficient construct validity
	SF-12	-	+	B	Indeterminate result rating for internal consistency
	SF-36	-	+	B	Indeterminate result rating for internal consistency
	WHOQOL-Bref	+	-	B	Indeterminate result rating for internal consistency
Life impact and neuromuscular function					
	JOACMEQ	+	+	B	
Neuromuscular function					
	10 s step test	+	-	C	High-quality evidence for insufficient criterion validity
	30MWT	+	-	C	High-quality evidence for insufficient responsiveness
	9-Hole peg test	++	-	B	
	BBS	++	-	B	
	European Myelopathy Scale	+	-	B	
	Foot tapping test	+	-	C	High-quality evidence for insufficient criterion validity
	Grip-and-release test	+	-	B	
	JOA	-	+	B	
	MDI	+	-	B	
	mJOA	-	+	C	High-quality evidence for insufficient criterion and construct validity
	Nurick scale	+	-	B	
	P-mJOA	+	-	B	
	Ranawat classification of disease severity	-	-	B	
	Triangle step test	+	-	B	
Pain and neuromuscular function					
	QuickDASH	-	-	B	
Pain					
	NDI	+	+	B	
	Arm pain score	-	+	B	
	Neck pain score	+	+	B	
	VAS for pain	+	+	B	
Radiology					
	Cobb's method	+	-	B	
	CT (Tsuayama's classification, 2D and 3D)	+	-	B	

Continued

Table 4 Continued

Domain	Instrument	Feasibility	Interpretability	Recommendation category	Recommendation justification
	CT (Tsuyama's classification, lateral + axial)	+	–	B	
	Isihara's cervical curvature index	+	–	B	
	MRI (depiction of intramedullary hyperintensity at eight cervical disc levels, T2W, 1.5-T or 3-T)	+	–	B	
	MRI (Kang's classification, 1.5-T or 3-T)	+	–	B	
	MRI (Muhle's classification, 1.5-T)	+	–	B	
	MRI (Vaccaro's classification, 1.5-T)	+	–	B	
	X-rays (computer-assisted measurement of length and thickness)	+	–	B	

Feasibility: ++=No barriers; +=Minimal barriers; –=Barriers

Interpretability: +=Interpretable; –= Uninterpretable, due to absence of anchor-based MCIDs ^{23–25}

Recommendation category: A=measurement instruments with evidence for sufficient content validity (any level) AND at least low-quality evidence for sufficient internal consistency; B=measurement instruments categorised not in A or C; C=measurement instruments with high-quality evidence for an insufficient measurement property.

BBS, Berg Balance Scale; EQ-5D, EuroQol-5 Dimension; JOA, Japanese Orthopaedic Association; JOACMEQ, Japanese Orthopaedic Association Cervical Myelopathy Evaluation Questionnaire; MDI, Myelopathy Disability Index; mJOA, modified Japanese Orthopaedic Association; 30MWT, 30-m Walking Test; NDI, Neck Disability Index; P-mJOA, patient-derived version of the mJOA; SF-12, 12-Item Short Form Health Survey; SF-36, 36-Item Short Form Health Survey; VAS, Visual Analogue Scale; WHOQOL-Bref, World Health Organisation Quality of Life.

sample sizes <100). Importantly, only one study formulated a hypothesis a priori.³⁵

Responsiveness

Sixteen measurement instruments were evaluated for responsiveness, including the JOACMEQ, JOA, mJOA, NDI, SF-12 and SF-36 (online supplemental table 12). The mJOA was the only score to attain high-quality evidence for sufficient responsiveness (effect size range: 0.87–1.0, total sample size: 352). The 30MWT, on the other hand, was the only score to attain high-quality evidence for insufficient responsiveness (standardised response mean: 0.3, total sample size: 484). The quality of the evidence of most of the remaining tools ranged from 'very low' to 'moderate' due to (1) imprecision (ie, sample sizes <100) and (b) uncertainty of the statistical methods.

Clinically important differences

Ten measurement instruments were evaluated for clinically important differences, including the JOACMEQ, JOA, mJOA, NDI, arm and neck pain scores, SF-12, SF-36 and VAS for pain (online supplemental table 3). From these, 7 of 10 attained a sufficient rating, including the JOACMEQ, JOA, mJOA, NDI and SF-36. Only anchor-based measures were accepted for the assessment of the MCID.^{23–25 36–39}

Interpretability and feasibility

Interpretability and feasibility were described using criteria adapted from the COSMIN methodology (online supplemental tables 3 and 4, respectively). Interpretability was summarised in terms of the degree to which

clinicians may assign qualitative meaning to the scores or change in scores (ie, the clinically important differences), while feasibility was described in terms of the ease of application of the measurement instrument. No or minimal application barriers were identified for most outcome measures (table 4). Nine instruments were, however, deemed uninterpretable due to the absence of anchor-based MCIDs.^{23–25}

Recommendations

No category A recommendations were made as no measurement instrument had sufficient evidence for content validity (table 4 and figure 2). Furthermore, five instruments were recommended for category C due to the availability of high-quality evidence for insufficient criterion validity, construct validity and/or responsiveness. Most instruments were classed into category B due to the notable absence of high-quality evidence for most measurement properties.

In light of these results, and given both (1) the very strict quality standards of the COSMIN framework and (2) that the absence of category A evidence is not the same as presence of poor-quality evidence, we propose that instruments most suitable for use should be interpretable by clinicians and offer qualitative meaning to either clinicians or people with lived experience of DCM (ie, they should have an available assessment of clinically important differences). To this end, the measurement properties of the nine interpretable instruments are presented in table 5: the arm and neck pain scores; SF-12 and SF-36; JOA, mJOA and JOACMEQ; NDI; and VAS

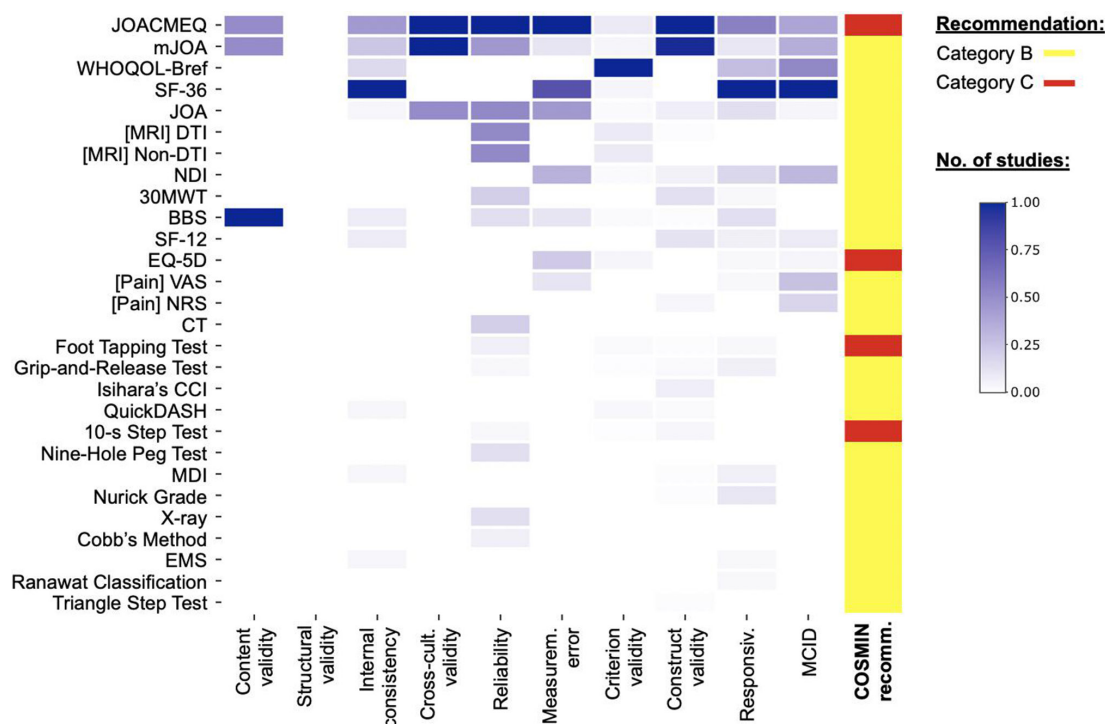


Figure 2 Number of studies for each outcome measure and property (normalised). Included studies reported on at least one of the 10 COSMIN properties for all instruments. No instrument had evidence for all 10 properties and <50% (13/29) of instruments had evidence for at least one property per measurement domain (see table 2 for definitions). Notably, no instruments were evaluated for structural validity, attained sufficient evidence for content validity or obtained a category A recommendation based on COSMIN criteria. 30MWT, 30-m Walking Test; BBS, Berg Balance Scale; COSMIN, Consensus-based Standards for the selection of health Measurement Instruments; EQ-5D, EuroQol-5 Dimension; JOA, Japanese Orthopaedic Association; JOACMEQ, Japanese Orthopaedic Association Cervical Myelopathy Evaluation Questionnaire; MDI, Myelopathy Disability Index; mJOA, modified Japanese Orthopaedic Association; NDI, Neck Disability Index; P-mJOA, patient-derived version of the mJOA; SF-12, 12-Item Short Form Health Survey; SF-36, 36-Item Short Form Health Survey; VAS, Visual Analogue Scale; WHOQOL-Bref, World Health Organisation Quality of Life

for pain. These include one score with insufficient criterion and construct validity (ie, mJOA) and six scores with barriers to application.

DISCUSSION

DCM is measured and reported inconsistently across clinical trials.¹⁻⁴ In light of these inconsistencies, AO Spine launched RECODE-DCM (www.aospine.org/recode) with the aim of creating a 'research toolkit' that helps to accelerate knowledge discovery and improve outcomes in DCM. One of the objectives of the RECODE-DCM initiative was to develop a CMS.^{3 6 26} This systematic review consists of an initial step towards building this CMS by identifying tools that have been used in DCM research and examining their quality, in accordance with the COSMIN standards.²³⁻²⁵

Overall, we identified 29 instruments with at least 1 in 10 measurement properties evaluated (figure 2); none, however, had evaluations for all 10 properties and <50% had more than one property evaluated per measurement domain (ie, reliability, validity and responsiveness) (table 2). We also noted a paucity in the quantity and quality of studies evaluating DCM instruments; this is visible by the absence of category A recommendations

and the classification of most tools in category B (table 4). Acknowledging both the stringency of the COSMIN standards and that absence of category A evidence is not equivalent to presence of poor-quality evidence, we proposed nine instruments that seem interpretable to clinicians and appear to offer qualitative meaning to clinicians and people with lived experience of DCM. These instruments are the SF-12 and SF-36; JOA, mJOA, and JOACMEQ; NDI; and VAS for pain (table 5).

The fact that most outcomes received B-category recommendations due to absence of high-quality evidence is not unexpected. In this review, the most common reasons for low-quality evidence, as per the COSMIN guidelines, were (1) important methodological flaws in study design or statistical methods, (2) uncertainty of approaches used to analyse the data and (3) imprecision due to sample size below the recommended power and significance levels. The rigour (or stringency) of the COSMIN standards may have accentuated these limitations due to the highly specific nature of some standards and the expectation of psychometric expertise within the DCM context. For example, results for internal consistency must be rated 'indeterminate' if there is not at least low-quality evidence for structural validity. No such studies were available in

Table 5 Interpretable measurement instruments

Domain	Instrument	Psychometric properties*	Feasibility	Recommendation category
Life impact				
	SF-12	Cronbach's α coefficient (0.77)	-	B
	MCS	SCB (51.5)		
	PCS	SCB (30.1) Responsiveness: SF-12 PCS (mean change score: 8.17)		
	SF-36	Cronbach's α coefficient (0.79–0.93) Responsiveness: SF-36 (normalised change: 0.32)	-	B
	MCS	MDC or SDC (distribution: 3.3–5.7) MCID (distribution: 3.4–6.8, anchor: 3.0–7.4) Construct validity: Arm pain score (Pearson's correlation: -0.23) mJOA scale (Pearson's correlation: 0.19) NDI (Spearman's rank correlation: -0.17) Neck pain score (Pearson's correlation: -0.28) SF-12 PCS (Pearson's correlation: 0.01) Responsiveness: SF-36 MCS (effect size range: 0.81, sensitivity: 0.67)		
	PCS	MDC or SDC (distribution: 5.2–5.7, anchor: 4.9) MCID (distribution: 2.9–5.5, distribution: 10, anchor: 3.9–9.6) SCB ¹⁶ Criterion validity (Likert scale): AUC: 0.67–0.69 Construct validity: Arm pain score (Pearson's correlation: -0.44) mJOA scale (Pearson's correlation: 0.43) NDI (Spearman's rank correlation: -0.49) Neck pain score (Pearson's correlation: -0.41) SF-12 PCS (Pearson's correlation: -0.29) Responsiveness: SF-36 PCS (effect size range: 0.84, sensitivity: 0.85)		
Life impact and neuromuscular function				
	JOACMEQ	Patient comprehensibility: 'No questions elicited no answer or 'I am not sure' in more than 5% of patients' Test-retest stability: Cronbach's α coefficient (0.91) Forward-backward translation (Persian and Thai): n/a	+	B
	Bladder function	Cronbach's α coefficient (0.32–0.74) Test-retest stability: ICC (0.62) MDC or SDC (distribution: 7.7) MCID (anchor: 6.0) Responsiveness: JOACMEQ bladder function (AUC: 0.82, effect size: 0.33, mean change score: 18.0)		
	Cervical spine function	Cronbach's α coefficient (0.77–0.78) Test-retest stability: ICC (0.63) MDC or SDC (distribution: 12.9, anchor: 12.5) MCID (anchor: 2.5) Criterion validity (Likert scale): AUC: 0.58 Responsiveness: JOACMEQ cervical spine function (AUC: 0.72, Effect size: 0.28, Mean change score: 25.8)		

Continued

Table 5 Continued

Domain	Instrument	Psychometric properties*	Feasibility	Recommendation category
	Lower extremity function	Cronbach's α coefficient (0.80–0.86) Test–retest stability: ICC (0.83) MDC or SDC (distribution: 6.6, anchor: 8.5) MCID (anchor: 8.5–9.5) Criterion validity (Likert scale): AUC: 0.66–0.70 Construct validity: NDI (Pearson's correlation: –0.66) SF-12 MCS (Spearman's rank correlation: 0.40) SF-12 PCS (Spearman's rank correlation: 0.29) Responsiveness: JOACMEQ quality of life (AUC: 0.83, effect size: 0.46, mean change score: 23.7)		
	QOL	Cronbach's α coefficient (0.80–0.86) Test–retest stability: ICC (0.83) MDC or SDC (distribution: 6.6, anchor: 8.5) MCID (anchor: 8.5–9.5) Criterion validity (Likert scale): AUC: 0.66–0.70 Construct validity: NDI (Pearson's correlation: –0.66) SF-12 MCS (Spearman's rank correlation: 0.40) SF-12 PCS (Spearman's rank correlation: 0.29) Responsiveness: JOACMEQ quality of life (AUC: 0.83, effect size: 0.46, mean change score: 23.7)		
	Upper extremity function	Cronbach's α coefficient (0.72–0.74) Test–retest stability: ICC (0.93) MDC or SDC (distribution: 9.5, anchor: 6.1) MCID (anchor: 2.5–13.0) Responsiveness: JOACMEQ upper extremity function (AUC: 0.74, effect size: 0.17, mean change score: 10.7)		
Neuromuscular function				
	JOA	Cronbach's α coefficient (0.72) Forward–backward translation (Brazilian Portuguese): Comprehension rate (>81.2%) Interobserver reliability: ICC (0.81) MDC or SDC (distribution: 1.0, anchor: 2.5) LOA (1.2(–1.2 to 3.6)) MCID (anchor: 2.5) Criterion validity (Likert scale): AUC: 0.59–0.62 Construct validity: JOACMEQ QOL (Spearman's rank correlation: 0.41) mJOA (Spearman's rank correlation: 0.87) NDI (Spearman's rank correlation: –0.50 to –0.76) SF-12 MCS (Spearman's rank correlation: –0.05) SF-12 PCS (Spearman's rank correlation: 0.50) Responsiveness: JOA (mean change score: 4.6, normalised change: 0.21) JOA motor function of lower extremity (mean change score: 0.60) mJOA (Spearman's rank correlation: 0.75)	–	B
	Bladder function	Intraobserver reliability ($\kappa=0.64$) Interobserver reliability ($\kappa=0.47$)		
	Motor function of fingers	Intraobserver reliability ($\kappa=0.68$) Interobserver reliability ($\kappa=0.53$)		

Continued

Table 5 Continued

Domain	Instrument	Psychometric properties*	Feasibility	Recommendation category
	Motor function of shoulder and elbow	Intraobserver reliability ($\kappa=0.50$) Interobserver reliability ($\kappa=0.31$)		
	Motor function of lower extremity	Intraobserver reliability ($\kappa=0.55$) Interobserver reliability ($\kappa=0.49$)		
	Sensory function of lower extremity	Intraobserver reliability ($\kappa=0.54$) Interobserver reliability ($\kappa=0.58$)		
	Sensory function of upper extremity	Intraobserver reliability ($\kappa=0.51$) Interobserver reliability ($\kappa=0.42$)		
	mJOA	Cronbach's α coefficient (0.60–0.63) Forward-backward translation (Brazilian Portuguese and Italian): n/a Test-retest stability (Spearman's rank correlation: 0.91) Intraobserver reliability (ICC: 0.87) Interobserver reliability (ICC: 0.97, $\kappa=0.80$) MDC or SDC (distribution: 2.1) MCID (distribution: 1.2–1.4, anchor: 1.3–3.1) SCB ¹⁴ Criterion validity (Nurick scale): Spearman's rank correlation: –0.41 Pearson's correlation: –0.62 to –0.63 Construct validity: 30MWT (Pearson's correlation: –0.38) EQ-5D (Spearman's rank correlation: 0.42) JOACMEQ QOL (Spearman's rank correlation: 0.41) NDI (Spearman's rank correlation: –0.51, Pearson's correlation: –0.33 to –0.34) SF-12 MCS (Pearson's correlation: 0.03) SF-12 PCS (Pearson's correlation: 0.42) SF-36 MCS (Pearson's correlation: 0.25) SF-36 PCS (Pearson's correlation: 0.30) Responsiveness: mJOA (effect size: 0.87–1.0, normalised change: 1.47)	–	C
	Motor dysfunction of lower extremities	Interobserver reliability (ICC: 0.73) Criterion validity (Nurick scale): Pearson's correlation: –0.65 to –0.68 Construct validity: 30MWT (Pearson's correlation: –0.43) NDI (Pearson's correlation: –0.31) SF-36 MCS (Pearson's correlation: 0.21) SF-36 PCS (Pearson's correlation: 0.31–0.50)		
	Motor dysfunction of upper extremities	Interobserver reliability (ICC: 0.77) Criterion validity (Nurick scale): Pearson's correlation: –0.42 Construct validity: 30MWT (Pearson's correlation: –0.21) NDI (Pearson's correlation: –0.24) SF-36 MCS (Pearson's correlation: 0.20) SF-36 PCS (Pearson's correlation: 0.22)		
	Sensory dysfunction of sphincter dysfunction	Interobserver reliability (ICC: 0.78) Criterion validity (Nurick scale): Pearson's correlation: –0.25 Construct validity: 30MWT (Pearson's correlation: –0.23) NDI (Pearson's correlation: –0.16) SF-36 MCS (Pearson's correlation: 0.08) SF-36 PCS (Pearson's correlation: 0.06)		

Continued



Table 5 Continued

Domain	Instrument	Psychometric properties*	Feasibility	Recommendation category
	Sensory dysfunction of upper extremities	Interobserver reliability (ICC: 0.93) Criterion validity (Nurick scale): Pearson's correlation: -0.23 Construct validity: 30MWT (Pearson's correlation: -0.05) NDI (Pearson's correlation: -0.23) SF-36 MCS (Pearson's correlation: 0.19) SF-36 PCS (Pearson's correlation: 0.19)		
Pain				
	NDI	MDC or SDC (distribution: 6.2%, anchor: 5.2%) MCID (anchor: 5-13) SCB (anchor: 9.5-36) Criterion validity (Likert scale): AUC: 0.66-0.75 Construct validity: Arm pain score (Pearson's correlation: 0.68) mJOA (Pearson's correlation: -0.36) Neck pain score (Pearson's correlation: 0.64) SF-12 MCS (Pearson's correlation: -0.40) SF-12 PCS (Pearson's correlation: -0.54) Responsiveness: Anchor (AUC: 0.66) NDI (mean change score: -15.8)	+	B
	Pain, 'Numeric rating scale' (arm pain score)	MCID (anchor: 2.5) SCB (3.5) Construct validity: mJOA (Pearson's correlation: -0.19) Neck pain score (Pearson's correlation: 0.72)	-	B
	Pain, 'Numeric rating scale' (neck pain scores)	MCID (anchor: 2.5) SCB (3.5) Construct validity: mJOA (Pearson's correlation: -0.07)	-	B
	VAS for pain	MDC or SDC (distribution: 3.1) MCID (distribution: 24.0-30.0, anchor: 0.4-2.7) SCB (1.1)	+	B

n/a=No info available

Feasibility: ++=No barriers; +=Minimal barriers; -=Barriers

Interpretability: +=Interpretable; -=Uninterpretable, due to absence of anchor-based MCIDs²³⁻²⁵

Recommendation category: A=measurement instruments with evidence for sufficient content validity (any level) AND at least low-quality evidence for sufficient internal consistency; B=Measurement instruments categorised not in A or C; C=measurement instruments with high-quality evidence for an insufficient measurement property.

*Comparators shown as indented tools

AUC, area under curve; EQ-5D, EuroQol-5 Dimension; JOA, Japanese Orthopaedic Association; JOACMEQ, Japanese Orthopaedic Association Cervical Myelopathy Evaluation Questionnaire; LOA, limits of agreement; MCID, minimal clinically important difference; MCS, mental component summary; MDC, minimal detectable change; mJOA, modified Japanese Orthopaedic Association; 30MWT, 30-m Walking Test; NDI, Neck Disability Index; PCS, physical component summary; SCB, substantial clinical benefit; SDC, smallest detectable change; SF-12, 12-Item Short Form Health Survey; SF-36, 36-Item Short Form Health Survey.

this review, possibly because this is a more recent and complex criterion, or because of the search or selection criteria. Similarly, studies on content validity cannot score higher than 'inadequate' if there are no recordings/verbatim transcriptions of patient focus groups or interviews. Likewise, analyses of reliability cannot score higher than 'doubtful' if statistics other than the Pearson or Spearman correlation coefficients are used. These thresholds of acceptability may account for some of the lacking information and are an important entry challenge for instruments into DCM research—a field where the routine involvement of stakeholders with lived experience is at an

early stage,^{3 8} inconsistent study reporting is prevalent,^{2 4} few studies have involved >100 patients, and where there is a bias in the availability of measurement literature (ie, some tools, such as the SF-12, are used because they are the only tools available and, therefore, have available literature due to their routine use). From the application of these COSMIN criteria in other research fields, however, it appears that these methodological deficiencies are not exclusive to DCM instruments, including those in current use.⁴⁰⁻⁴² The lack of high-quality assessments, thus, should not necessarily imply that (1) the identified

outcome measures are generally inadequate, or (b) that the COSMIN standards are not fit for the DCM context.

Measurement rigour is universally important and, in DCM, particularly relevant as the development of new instruments is a top 10 research priority. This rank reinforces the decision of the steering committee to make the initial CMS recommendations based existing on tools, rather than on tools under development.²⁶ This decision was taken recognising that the success of a CMS requires widespread adoption, and that the adoption of clinical recommendations can be challenging without stakeholder awareness, familiarity and/or confidence.^{43–46} We hypothesised that asking the global field to align with new innovations would be more challenging, and premature, at this stage. Thus, for this first iteration of the DCM CMS, there is a focus on current instruments in academic usage. While, currently, few have met the bar set by the COSMIN methodology, there are nine reasonable candidates using our post-hoc thresholds (table 5). Ultimately, the CMS process will need to lean significantly on the expertise of those involved in the consensus phase in order to make final recommendations that are methodologically rigorous and representative of those with lived experience.

Despite its conscientious design, this systematic review has limitations. In searching for existing instruments, we have neither identified nor assessed tools under development, or those currently being translated into clinical or research settings or published in languages other than English. To the extent that DCM instruments are currently in use, however, this review only identified tools in four of the six core domains from RECODE-DCM's minimum dataset,³³ and did not consider the construct of the disease as a factor in evaluating the outcomes. For those missing outcomes, focused scoping reviews (informed by a gap analysis that will be published separately) will be conducted in the future. Next, clinician-reported outcome measures and performance-based outcome measures were analysed with the exact same methods as PROMs. While COSMIN explicitly allows this,^{23–25} methods may be differentially adapted to tailor to these distinct instrument types; we chose not to do so out of prudence and consistency, and results across these instrument groups should be interpreted accordingly. Feasibility and interpretability were also evaluated using bespoke criteria which, despite being adapted from the COSMIN methodology, may not weigh all criteria accurately. Importantly, our decision to shortlist the clinically interpretable instruments was made a posteriori due to the unexpected absence of category A recommendations. This decision was informed by our judgement that instruments in a CMS should be interpretable by clinicians and offer qualitative meaning to clinicians and people with lived experience. While the COSMIN taxonomy does indeed class interpretability as an important and standalone characteristic,^{23–25} the aforementioned shortlist may inevitably represent a placement bias. Notably, some nuances of different versions of measurement instruments

(eg, mJOA) were not extensively evaluated.⁴⁷ Lastly, and as is frequently the case in this body of reviews,^{40–42} none of the authors is specifically trained in measurement theory and, therefore, this work represents our best attempt to implement the guidelines and standards set forward by the COSMIN methodology in the context of DCM.

CONCLUSIONS

Currently, none of the measurement instruments used in DCM holds sufficient evidence to meet the COSMIN criteria for a strong recommendation for use. However, there are leading contenders that appear to offer qualitative meaning to clinicians and people with lived experience of DCM; namely, the SF-12 and SF-36; JOA, mJOA, and JOACMEQ; NDI; and VAS for pain. The findings of this review will inform a consensus process to form a CMS for DCM. As the development of new assessments for DCM is an active research priority, greater awareness of the COSMIN framework is pertinent to DCM researchers.

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SUPPLEMENTARY INFORMATION**Supplementary Table 1. Full search phrases used for MEDLINE and EMBASE on 4 August 2020**

Ovid MEDLINE	2389 articles
Cervical spine concept	
1	exp Cervical Vertebrae/ or exp Cervical Cord/ or cervical.tw
DCM concept	
2	Exp Spinal Cord Diseases/ or Exp Spinal Diseases/
3	degenerat*.tw
4	2 and 3
5	Myelopath*.tw
6	Myeloradiculopath*.tw
7	Radiculopath*.tw
8	Exp Spinal Cord Compression/
9	Exp "Ossification of the Posterior Longitudinal Ligament"/
10	Ossification of the Posterior Longitudinal Ligament.tw
11	OPLL.tw
12	Exp Spinal Stenosis/
13	Cervical stenosis.tw
14	Exp Spondylosis/
15	Spondylosis.tw
16	Spondylotic.tw
17	Degenerative cervical myelopathy.tw
18	DCM.tw
19	Cervical spondylotic myelopathy.tw
20	CSM.tw
21	4 or 5 or 6 or 7 or 8 or 9 or 10 or 11 or 12 or 13 or 14 or 15 or 16 or 17 or 18 or 19 or 20
22	1 and 21
Tools for function concept (neurological/gait/mobility centric) concept	
23	Japanese Orthopaedic Association score.tw
24	JOA.tw
25	modified JOA.tw
26	mJOA.tw
27	Graded Redefined Assessment of Sensation Strength and Prehension.tw
28	GRASSP.tw
29	Quick Disability of the Arm Shoulder and Hand.tw
30	QuickDASH.tw

- 31 Myelopathy Disability Index.tw
- 32 MDI.tw
- 33 Nurick score.tw
- 34 Neck functional disability scale.tw
- 35 NFDS.tw
- 36 Neck Disability Index.tw
- 37 NDI.tw
- 38 Cooper myelopathy scale.tw
- 39 CMS.tw
- 40 European myelopathy score.tw
- 41 EMS.tw
- 42 Bournemouth questionnaire.tw
- 43 BQ.tw
- 44 Cervical spine outcomes questionnaire.tw
- 45 CSOQ.tw
- 46 Patient specific functional scale.tw
- 47 PSFS.tw
- 48 World Health Organization Quality of Life Instruments.tw
- 49 WHOQOL.tw
- 50 Grip and release test.tw
- 51 GRT.tw
- 52 Grip Dynamometer.tw
- 53 Triangle step test.tw
- 54 Foot tapping test.tw
- 55 30 m walking test.tw
- 56 30MWT.tw
- 57 10 m walking test.tw
- 58 10MWT.tw
- 59 Berg Balance Scale.tw
- 60 BBS.tw
- 61 GAITRite.tw
- 62 10 second step test.tw
- 63 9 hole peg test.tw
- 64 Prolo.tw
- 65 Mental component score.tw
- 66 MCS.tw
- 67 Physical component score.tw
- 68 PCS.tw

- 69 Hospital anxiety depression scale.tw
 70 HADS.tw
 71 Global rating of change.tw
 72 GROCC.tw
 73 23 or 24 or 25 or 26 or 27 or 28 or 29 or 30 or 31 or 32 or 33 or 34 or 35 or 36 or 37 or 38 or 39 or 40 or 41 or 42 or 43 or 44 or 45 or 46 or 47 or 48 or 49 or 50 or 51 or 52 or 53 or 54 or 55 or 56 or 57 or 58 or 59 or 60 or 61 or 62 or 63 or 64 or 65 or 66 or 67 or 68 or 69 or 70 or 71 or 72

Tools for QOL concept (including pain) concept

- 74 Exp "Quality of Life"/ or exp "Surveys and Questionnaires"/
 75 Short Form Health Survey.tw
 76 SF-36.tw
 77 SF-12.tw
 78 EQ-5D.tw
 79 Japanese Orthopaedic Association Cervical Myelopathy Evaluation Questionnaire.tw
 80 JOACMEQ.tw
 81 Visual Analogue Scale.tw
 82 VAS.tw
 83 Likert scale.tw
 84 Numeric pain rating scale.tw
 85 NPRS.tw
 86 North American Spine Satisfaction.tw
 87 NASS.tw
 88 74 or 75 or 76 or 77 or 78 or 79 or 80 or 81 or 82 or 83 or 84 or 85 or 86 or 87

Psychometric concept

- 89 Exp Psychometrics/
 90 Pyschometr*.tw
 91 (cliniometr* or clinometr*).tw.
 92 Outcome assessment*.tw
 93 exp Health Status Indicators/
 94 Exp "Reproducibility of Results"/
 95 Reproducib*.tw
 96 Exp Validation Study/
 97 exp Discriminant Analysis/
 98 (reliab* or unreliab* or valid* or coefficient or homogeneity or homogeneous or internal consistency).tw
 99 (cronbach* and (alpha or alphas)).tw.
 100 (item and (correlation* or selection* or reduction*)).tw

- 101 (agreement or precision or imprecision or precise values or test–retest).tw
- 102 (reliab* and (test or retest)).tw
- 103 (stability or interrater or inter-rater or intrarater or intra-rater or intertester or inter-tester or intratester or intra-tester or interobserver or inter-observer or intraobserver or intraobserver or intertechnician or inter-technician or intratechnician or intra-technician or interexaminer or inter-examiner or intraexaminer or intra-examiner or interassay or interassay or intraassay or intra-assay or interindividual or inter-individual or intraindividual or intra-individual or interparticipant or inter-participant or intraparticipant or intra-participant or kappa or kappas or repeatab*).tw
- 104 ((replicab* or repeated) and (measure or measures or findings or result or results or test or tests)).tw
- 105 (generaliza* or generalisa* or concordance).tw
- 106 (intraclass and correlation*).tw
- 107 Exp Observer Variation/
- 108 Observer variation.tw
- 109 (multitrait and scaling and (analysis or analyses)).tw
- 110 Measurement error*.tw
- 111 (item discriminant or interscale correlation* or error or errors or individual variability).tw
- 112 (variability and (analysis or values)).tw
- 113 (uncertainty and (measurement or measuring)).tw
- 114 Exp Diagnostic Errors/
- 115 Exp Data accuracy/
- 116 Exp Dimensional Measurement Accuracy/
- 117 Accuracy.tw
- 118 ((minimal or minimally or clinical or clinically) and (important or significant or detectable) and (change or difference)).tw
- 119 Minimally clinically important difference*.tw
- 120 MCID.tw
- 121 (small* and (real or detectable) and (change or difference)).tw
- 122 (meaningful change or ceiling effect or floor effect or Item response model or IRT or Rasch or Differential item functioning or DIF or computer adaptive testing or item bank or cross-cultural equivalence).tw
- 123 Exp Bias/ or exp Selection Bias/
- 124 Bias.tw
- 125 Exp “Predictive Value of Test”/

126 89 or 90 or 91 or 92 or 93 or 94 or 95 or 96 or 97 or 98 or 99 or 100 or 101 or 102 or 103 or 104 or 105 or 106 or 107 or 108 or 109 or 110 or 111 or 112 or 113 or 114 or 115 or 116 or 117 or 118 or 119 or 120 or 121 or 122 or 123 or 124 or 125

Combined concepts

127 73 or 88

128 22 and 126 and 127

EMBASE

1550 articles

Cervical spine concept

1 exp Cervical Vertebra/ or cervical spine/ or exp Cervical spinal cord/ or cervical.tw

DCM concept

2 Exp Spinal Cord Disease/ or Exp Spine Disease/

3 Exp degeneration/

4 degenerat*.tw

5 3 or 4

6 2 and 5

7 Myelopath*.tw

8 Myeloradiculopath*.tw

9 Exp radiculopathy/

10 Radiculopath*.tw

11 Exp Spinal Cord Compression/

12 Exp Posterior Longitudinal Ligament/ and exp ossification/

13 Ossification of the Posterior Longitudinal Ligament.tw

14 OPLL.tw

15 Exp vertebral canal stenosis/

16 Cervical stenosis.tw

17 Exp Cervical Spondylosis/

18 Exp Spondylosis/

19 Spondylosis.tw

20 Spondylotic.tw

21 Exp Cervical myelopathy/

22 Degenerative cervical myelopathy.tw

23 DCM.tw

24 Exp Cervical spondylotic myelopathy/

25 Cervical spondylotic myelopathy.tw

26 CSM.tw

27 6 or 7 or 8 or 9 or 10 or 11 or 12 or 13 or 14 or 15 or 16 or 17 or 18 or 19 or 20 or 21 or 22 or 23 or 24 or 25 or 26

28 1 and 27

Tools for function concept (neurological/gait/mobility centric) concept

- 29 Exp Japanese Orthopaedic Association score/
- 30 Japanese Orthopaedic Association score.tw
- 31 JOA.tw
- 32 modified JOA.tw
- 33 mJOA.tw
- 34 Exp "Disabilities of the Arm, Shoulder and Hand (score)"/
- 35 Graded Redefined Assessment of Sensation Strength and Prehension.tw
- 36 GRASSP.tw
- 37 Quick Disability of the Arm Shoulder and Hand.tw
- 38 QuickDASH.tw
- 39 Myelopathy Disability Index.tw
- 40 MDI.tw
- 41 Exp "Nurick (grade)"/
- 42 Nurick score.tw
- 43 Neck functional disability scale.tw
- 44 NFDS.tw
- 45 Exp Neck Disability Index/
- 46 Neck Disability Index.tw
- 47 NDI.tw
- 48 Cooper myelopathy scale.tw
- 49 CMS.tw
- 50 European myelopathy score.tw
- 51 EMS.tw
- 52 Bournemouth questionnaire.tw
- 53 BQ.tw
- 54 Cervical spine outcomes questionnaire.tw
- 55 CSOQ.tw
- 56 Patient specific functional scale.tw
- 57 PSFS.tw
- 58 World Health Organization Quality of Life Instruments.tw
- 59 WHOQOL.tw
- 60 Grip and release test.tw
- 61 GRT.tw
- 62 Grip Dynamometer.tw
- 63 Triangle step test.tw
- 64 Foot tapping test.tw

- 65 30 m walking test.tw
- 66 30MWT.tw
- 67 10 m walking test.tw
- 68 10MWT.tw
- 69 Berg Balance Scale.tw
- 70 BBS.tw
- 71 GAITRite.tw
- 72 10 second step test.tw
- 73 9 hole peg test.tw
- 74 Prolo.tw
- 75 Mental component score.tw
- 76 MCS.tw
- 77 Physical component score.tw
- 78 PCS.tw
- 79 Hospital anxiety depression scale.tw
- 80 HADS.tw
- 81 Global rating of change.tw
- 82 GROC.tw
- 83 29 or 30 or 31 or 32 or 33 or 34 or 35 or 36 or 37 or 38 or 39 or 40 or 41 or 42 or 43 or 44 or 45 or 46 or 47 or 48 or 49 or 50 or 51 or 52 or 53 or 54 or 55 or 56 or 57 or 58 or 59 or 60 or 61 or 62 or 63 or 64 or 65 or 66 or 67 or 68 or 69 or 70 or 71 or 72 or 73 or 74 or 75 or 76 or 77 or 78 or 79 or 80 or 81 or 82

Tools for QOL concept (including pain) concept

- 84 Short Form Health Survey.tw
- 85 Exp Short Form 36/
- 86 SF-36.tw
- 87 Exp Short Form 12/
- 88 SF-12.tw
- 89 Exp "European Quality of Life 5 Dimensions questionnaire"/
- 90 EQ-5D.tw
- 91 Exp Japanese Orthopaedic Association Cervical Myelopathy Evaluation/
- 92 Japanese Orthopaedic Association Cervical Myelopathy Evaluation Questionnaire.tw
- 93 JOACMEQ.tw
- 94 Exp visual analog scale/
- 95 Visual Analogue Scale.tw
- 96 VAS.tw
- 97 Exp Likert scale/
- 98 Likert scale.tw

- 99 Numeric pain rating scale.tw
 100 NPRS.tw
 101 North American Spine Satisfaction.tw
 102 NASS.tw
 103 84 or 85 or 86 or 87 or 88 or 89 or 90 or 91 or 92 or 93 or 94 or 95 or 96 or 97 or 98 or 99
 or 100 or 101 or 102

Psychometric concept

- 104 Exp Psychometry/
 105 Pyschometr*.tw
 106 (clinimetr* or clinometr*).tw.
 107 Outcome assessment*.tw
 108 exp Health Status Indicator/
 109 Exp Reproducibility/
 110 Reproducib*.tw
 111 Exp Validation Study/
 112 exp Discriminant Analysis/
 113 (reliab* or unreliab* or valid* or coefficient or homogeneity or homogeneous or internal
 consistency).tw
 114 (cronbach* and (alpha or alphas)).tw.
 115 (item and (correlation* or selection* or reduction*)).tw
 116 (agreement or precision or imprecision or precise values or test–retest).tw
 117 (reliab* and (test or retest)).tw
 118 (stability or interrater or inter-rater or intrarater or intra-rater or intertester or inter-tester or
 intratester or intra-tester or interobserver or inter-observer or intraobserver or
 intraobserver or intertechnician or inter-technician or intratechnician or intra-technician or
 interexaminer or inter-examiner or intraexaminer or intra-examiner or interassay or
 interassay or intraassay or intra-assay or interindividual or inter-individual or
 intraindividual or intra-individual or interparticipant or inter-participant or intraparticipant or
 intra-participant or kappa or kappas or repeatab*).tw
 119 ((replicab* or repeated) and (measure or measures or findings or result or results or test
 or tests)).tw
 120 (generaliza* or generalisa* or concordance).tw
 121 (intraclass and correlation*).tw
 122 Exp Observer Variation/
 123 Observer variation.tw
 124 (multitrait and scaling and (analysis or analyses)).tw
 125 Measurement error*.tw
 126 (item discriminant or interscale correlation* or error or errors or individual variability).tw

- 127 (variability and (analysis or values)).tw
128 (uncertainty and (measurement or measuring)).tw
129 Exp Diagnostic Error/
130 Exp Data accuracy/
131 Exp Dimensional Measurement Accuracy/
132 ((minimal or minimally or clinical or clinically) and (important or significant or detectable) and (change or difference)).tw
133 Minimally clinically important difference*.tw
134 MCID.tw
135 (small* and (real or detectable) and (change or difference)).tw
136 (meaningful change or ceiling effect or floor effect or Item response model or IRT or Rasch or Differential item functioning or DIF or computer adaptive testing or item bank or cross-cultural equivalence).tw
137 Exp Bias/ or exp Selection Bias/
138 Bias.tw
139 104 or 105 or 106 or 107 or 108 or 109 or 110 or 111 or 112 or 113 or 114 or 115 or 116 or 117 or 118 or 119 or 120 or 121 or 122 or 123 or 124 or 125 or 126 or 127 or 128 or 129 or 130 or 131 or 132 or 133 or 134 or 135 or 136 or 137 or 138

Combined concepts

- 140 83 or 103
141 28 and 139 and 140
-

Supplementary Table 2. Study characteristics

Study	Country	Sample size	Psychometric properties	Outcome measures
Auffinger, Lall (1)	United States	30	MCID/SCB Measurement error	NDI VAS for pain SF-36
Augusto, Diniz (2)	Brazil	30	Cross-cultural validity/Measurement invariance Reliability Responsiveness	JOA NDI
Azimi, Rezaei (3)	Iran	87	Cross-cultural validity/Measurement invariance Responsiveness	JOACMEQ
Badhiwala, Witiw (4)	Canada	606	MCID/SCB	SF-36 mJOA Nurick Scale NDI
Bohm, Fehlings (5)	Multicenter/ Global	601	Reliability Hypotheses testing for construct validity Responsiveness	Walking tests (timed or steps) mJOA Nurick Scale NDI SF-36
Carreon, Glassman (6)	United States	505	MCID/SCB	NDI SF-36 "Numeric rating scale" for pain
Chang, Kong (7)	Korea	108	Reliability	CT / CTM
Chiba, Kato (8)	Japan		Reliability	X-rays
Chien, Lai (9)	Taiwan	45	Responsiveness MCID/SCB	JOACMEQ NDI
Chiu and Pang (10)	Hong Kong	72	Internal consistency Reliability Content validity Hypotheses testing for construct validity Criterion validity Measurement error	BBS mJOA

			Responsiveness	
Fukui, Chiba (11)	Japan	368	Content validity	JOACMEQ
Fukui, Chiba (12)	Japan	201	Reliability	JOACMEQ
Goyal, Murphy (13)	United States	118	Responsiveness	NDI SF-12
Gwinn, Iannotti (14)	United States	20	Reliability	X-rays Cobb's method
Hosono, Sakaura (15)	Japan	30	Reliability Criterion validity	Grip-and-release test JOA
Hosono, Takenaka (16)	Japan	48	Responsiveness	Grip-and-release test JOA
Kang, Lee (17)	Korea	82	Reliability	MRI (not DTI)
Kato, Oshima (18)	Japan	92	Measurement error Hypotheses testing for construct validity Responsiveness	JOA mJOA JOACMEQ NDI SF-12
Kato, Oshima (19)	Japan	101	Measurement error Criterion validity MCID/SCB	JOA Likert scale
Kato, Oshima (20)	Japan	101	Measurement error Criterion validity MCID/SCB	JOACMEQ NDI EQ-5D SF-36 Likert scale
King and Roberts (21)	United States	88	Internal consistency	SF-36
Ko, Choi (22)	Korea	357	Reliability	MRI (not DTI)
Kopjar, Tetreault (23)	USA	277	Responsiveness Hypotheses testing for construct validity Internal consistency	mJOA Nurick Scale NDI SF-36 Walking tests (timed or steps)
Latimer, Haden (24)	England	70	Responsiveness	SF-36 NDI

				VAS for pain MDI
Longo, Berton (25)	Italy	75	Cross-cultural validity/Measurement invariance Reliability Internal consistency Hypotheses testing for construct validity Responsiveness Criterion validity	mJOA Nurick Scale NDI SF-36
Lubelski, Alvin (26)	United States	119	Hypotheses testing for construct validity Responsiveness Criterion validity	mJOA Nurick Scale EQ-5D
Mihara, Kondo (27)	Japan	270	Hypotheses testing for construct validity	Grip-and-release test Triangle step test
Nakamoto, Oshima (28)	Japan	94	Internal consistency Hypotheses testing for construct validity Criterion validity	QuickDASH JOA NDI SF-36 "Numeric rating scale" for pain
Nakashima, Yukawa (29)	Japan	101	Hypotheses testing for construct validity	
Nicholson, Millhouse (30)	United States	235	Hypotheses testing for construct validity	MRI (not DTI) mJOA NDI SF-12 "Numeric rating scale" for pain Isihara's Cervical Curvature Index
Nikaido, Kikuchi (31)	Japan	87	Hypotheses testing for construct validity)	JOACMEQ SF-36
Numasawa, Ono (32)	Japan	126	Hypotheses testing for construct validity Responsiveness Reliability	JOA Foot tapping test Grip-and-release test
Olindo, Signate (33)	France	40	Reliability	9-Hole peg test MRI (not DTI) Nurick Scale mJOA Walking tests (timed or steps)

Park, Kim (34)	Korea	100	Reliability	MRI (not DTI)
Pratali, Smith (35)	Brazil		Cross-cultural validity	mJOA
Pratali, Smith (36)	Brazil	55	Reliability	mJOA
Rhee, Shi (37)	United States	100	Criterion validity Reliability Content validity	mJOA
Sato, Horikoshi (38)	Japan	66	Hypotheses testing for construct validity	MRI (DTI) JOA
Shim, Lee (39)	Korea	79	Reliability Criterion validity	MRI (not DTI)
Singh and Crockard (40)	England	100	Internal consistency Responsiveness	Odom's Criteria Nurick Scale Ranawat classification of disease severity MDI JOA EMS SF-36
Singh and Crockard (41)	United Kingdom	41	Hypotheses testing for construct validity	Walking tests (timed or steps) MDI Nurick Scale
Singh, Gnanalingham (42)	England	105	Internal consistency Criterion validity Responsiveness	SF-12 SF-36
Spurgas, Abbas (43)	USA	35	MCID/SCB	VAS for pain NDI SF-12 mJOA
Tetreault, Nouri (44)	Canada	755	MCID/SCB	mJOA NDI
Thakar and Rajshekhar (45)	India	51	MCID/SCB Responsiveness	VAS for pain Nurick Scale SF-36
Thakar, Christopher (46)	India	70	Internal consistency Criterion validity	WHOQOL-Bref SF-36

			Responsiveness MCID/SCB	Nurick Scale
Wada, Fukui (47)	Japan	137	Responsiveness	JOACMEQ JOA 10-s step test
Witayakom, Paholpak (48)	Thailand	70	Cross-cultural validity/Measurement invariance Reliability Internal consistency Hypotheses testing for construct validity	JOACMEQ SF-36
Yonenobu, Abumi (49)	Japan	29	Reliability	JOA
Yukawa, Kato (50)	Japan	163	Hypotheses testing for construct validity Reliability Criterion validity	10-s step test JOA Grip-and-release test
Zhang, Zhou (51)	China	142	Internal consistency Responsiveness MCID/SCB	SF-36 mJOA
Zhou, Zhang (52)	China	113	MCID/SCB Measurement error	mJOA SF-36

Supplementary Table 3. Interpretability (i.e., MCID and SCB).

Instrument	Result summary	Overall rating
EQ-5D	MCID: 0.05; total sample size: 101	Sufficient
JOA	MCID: 2.5; total sample size: 101	Sufficient
JOACMEQ		
Bladder function	MCID: 6.0; total sample size: 78	Sufficient
Cervical spine function	MCID: 2.5; total sample size: 179	Sufficient
Lower extremity function	MCID range 2.5–9.4; total sample size: 179	Sufficient
QOL	MCID range 8.5–9.5; total sample size: 179	Sufficient
Upper extremity function	MCID range 2.5–13.0; total sample size: 179	Sufficient
mJOA	MCID range 1.3–3.1; total sample size: 868	Sufficient
	SCB: 14; total sample size: 35	Indeterminate
NDI	MCID range 5–13; total sample size: 108	Sufficient
	SCB range 9.5–36; total sample size: 65	Indeterminate
Pain, "Numeric rating scale" (Arm pain)	MCID: 2.5; total sample size: 30	Indeterminate
	SCB: 3.5; total sample size: 30	
Pain, "Numeric rating scale" (Neck pain)	MCID: 2.5; total sample size: 30	Indeterminate
	SCB: 3.5; total sample size: 30	
SF-12		

MCS	SCB: 51.5; total sample size: 35	Indeterminate
PCS	SCB: 30.1; total sample size: 35	Indeterminate
SF-36		
MCS	MCID range 3.0–7.4; total sample size: 749	Sufficient
PCS	"MCID range 3.9–9.6; total sample size: 890 SCB: 16; total sample size: 30"	Sufficient
VAS for pain	MCID range 0.4–2.7; total sample size: 30	Sufficient
	SCB: 1.1; total sample size: 30	Indeterminate

Supplementary Table 4. Feasibility assessment.

Tool	Time (min)	Equipment	Training	License	Money	Ease of administration	Overall assessment
10-s step test	5	Minimal	No	No	No	Minimal barriers	Minimal barriers
30MWT	5	Minimal	No	No	No	Minimal barriers	Minimal barriers
9-Hole peg test	5	Yes	No	No	No	Barriers	Barriers
Berg Balance Scale	>15	Yes	Yes	No	No	Barriers	Barriers
Cobb's method (C2-C7)	5	Minimal	No	No	No	Minimal barriers	Minimal barriers
CT (Tsuyama's classification, 2D & 3D)	5	Minimal	No	No	No	Minimal barriers	Minimal barriers
CT (Tsuyama's classification, lateral + axial)	5	Minimal	No	No	No	Minimal barriers	Minimal barriers
EQ-5D European	5	Minimal	No	Yes	Yes	Minimal barriers	Minimal barriers
Myelopathy Scale	5	Minimal	No	No	No	Minimal barriers	Minimal barriers
Foot tapping test	5	Minimal	No	No	No	Minimal barriers	Minimal barriers
Grip-and-release test	5	Minimal	No	No	No	Minimal barriers	Minimal barriers
Isihara's Cervical Curvature Index	5	Minimal	No	No	No	Minimal barriers	Minimal barriers
JOA	5	No	No	No	No	No barriers	No barriers

JOACMEQ	5–15	Minimal	No	No	No	Minimal barriers	Minimal barriers
MDI	5	Minimal	No	No	No	Minimal barriers	Minimal barriers
mJOA	5	No	No	No	No	No barriers	No barriers
MRI (Depiction of intramedullary hyperintensity at eight cervical disc levels, T2W, 1.5-T or 3-T)	5–15	Minimal	No	No	No	Minimal barriers	Minimal barriers
MRI (Kang's classification, 1.5-T or 3-T)	5	Minimal	No	No	No	Minimal barriers	Minimal barriers
MRI (Muhle's classification, 1.5-T)	5	Minimal	No	No	No	Minimal barriers	Minimal barriers
MRI (Vaccaro's classification, 1.5-T)	5	Minimal	No	No	No	Minimal barriers	Minimal barriers
NDI	5	Minimal	No	No	No	Minimal barriers	Minimal barriers
Nurick scale	5	Minimal	No	No	No	Minimal barriers	Minimal barriers
P-mJOA	5	Minimal	No	No	No	Minimal barriers	Minimal barriers

Arm pain score	5	No	No	No	No	No barriers	No barriers
Neck pain score	5	No	No	No	No	No barriers	No barriers
QuickDASH	5	Minimal	No	Yes	Yes	Minimal barriers	Barriers
Ranawat classification of disease severity	5	No	No	No	No	No barriers	No barriers
SF-12	5	Minimal	No	Yes	Yes	Minimal barriers	Barriers
SF-36	5–15	Minimal	No	Yes	Yes	Minimal barriers	Barriers
Triangle step test	5	Minimal	No	No	No	Minimal barriers	Minimal barriers
VAS for pain	5	Minimal	No	No	No	Minimal barriers	Minimal barriers
WHOQOL-Bref	5–15	Minimal	No	No	No	Minimal barriers	Minimal barriers
X-rays (Computer-assisted measurement of length & thickness)	5–15	Minimal	No	No	No	Minimal barriers	Minimal barriers

Supplementary Table 5. Content validity.

Instrument	Result summary	Overall rating	Quality of evidence
BBS	Patient comprehensibility: Item discrimination index >0.589	Indeterminate	Very low
JOACMEQ	Patient comprehensibility: "No questions elicited no answer or "I am not sure" in more than 5% of patients"	Indeterminate	Very low
P-mJOA	Patient comprehensibility: "In patients preferring to complete the mJOA them- selves, the most popular answers were: "ease of answering the questions" (n = 33), "understanding of the questions" (n = 17)"	Indeterminate	Very low

Supplementary Table 6. Internal consistency.

Instrument	Result summary	Overall rating	Quality of evidence
BBS	Cronbach's alpha range 0.95–0.98; consistent; total sample size: 72	Indeterminate	Moderate
European Myelopathy Scale	Cronbach's alpha: 0.68; consistent; total sample size: 100	Indeterminate	Low
JOA	Cronbach's alpha: 0.72; consistent; total sample size: 100	Indeterminate	Low
JOACMEQ	Cronbach's alpha: 0.91; total sample size: 70	Indeterminate	Moderate
Bladder function	Cronbach's alpha range 0.77–0.78; consistent; total sample size: 157	Indeterminate	High
Cervical spine function	Cronbach's alpha range 0.75–0.88; consistent; total sample size: 157	Indeterminate	High
QOL	Cronbach's alpha range 0.80–0.86; consistent; total sample size: 157	Indeterminate	High
Upper extremity function	Cronbach's alpha range 0.72–0.74; consistent; total sample size: 157	Indeterminate	High
MDI	Cronbach's alpha: 0.92; consistent; total sample size: 100	Indeterminate	Low
mJOA	Cronbach's alpha range 0.60–0.63; consistent; total sample size: 352	Indeterminate	High
QuickDASH	Cronbach's alpha: 0.94; consistent; total sample size: 94	Indeterminate	Very low
SF-12	Cronbach's alpha: 0.77; consistent; total sample size: 105	Indeterminate	n/a

SF-36	Cronbach's alpha range 0.79–0.93; consistent; total sample size: 473	Indeterminate	n/a
WHOQOL-Bref	Cronbach's alpha range 0.86–0.87; consistent; total sample size: 38	Indeterminate	n/a

n/a = No info available

Supplementary Table 7. Cross-cultural validity.

Instrument	Result summary	Overall rating	Quality of evidence
JOA	Forward-backward translation [Brazilian Portuguese] Comprehension rate: >81.2%	Indeterminate	Very low
JOACMEQ	Forward-backward translation [Persian and Thai] No info available	Indeterminate	Very low
mJOA	Forward-backward translation [Brazilian Portuguese and Italian] No info available	Indeterminate	Very low

Supplementary Table 8. Reliability.

Instrument	Result summary	Overall rating	Quality of evidence
10-s step test	Test-retest stability: Spearman's rank correlation: 0.89; total sample size: 163	Indeterminate	Low
30MWT	Test-retest stability: Pearson's correlation range 0.89–1.00; total sample size: 16	Indeterminate	Very low
9-Hole peg test	Intra-observer reliability: ICC range 0.97–0.98; consistent; total sample size: 41 Inter-observer reliability: ICC range 0.97–0.99; consistent; total sample size: 41	Sufficient	Very low
BBS*	Test-retest stability: ICC: 0.99; total sample size: 32 Inter-observer reliability: ICC: 0.99; total sample size: 32	Sufficient	Very low
	Test-retest stability: Kappa: 0.67; total sample size: 32 Inter-observer reliability: Kappa: 0.43; total sample size: 32	Insufficient	Very low
Cobb's method	Intra-observer reliability: ICC: 0.84; total sample size: 20 Inter-observer reliability: ICC: 0.77; total sample size: 20	Sufficient	Very low
CT	Intra-observer reliability:	Sufficient	Moderate

(Tsuyama's classification, 2D & 3D)	Kappa range 0.85–0.86; consistent; total sample size: 108 Inter-observer reliability: Kappa range 0.71–0.76; consistent; total sample size: 108		
CT (Tsuyama's classification, lateral + axial)	Intra-observer reliability: Kappa: 0.67; total sample size: 108 Inter-observer reliability: Kappa: 0.51; total sample size: 108	Insufficient	Moderate
Foot tapping test	Test–retest stability: Pearson's correlation range 0.90–0.93; total sample size: 126	Indeterminate	Low
Grip-and-release test	Inter-observer reliability: ICC: 0.99; total sample size: 30	Sufficient	Very low
JOA	Inter-observer reliability: ICC: 0.81; total sample size: 29	Sufficient	Very low
Bladder function	Intra-observer reliability: Kappa: 0.64; total sample size: 29 Inter-observer reliability: Kappa: 0.47; total sample size: 29	Insufficient	Very low
Motor function of fingers	Intra-observer reliability: Kappa: 0.68; total sample size: 29 Inter-observer reliability: Kappa: 0.53; total sample size: 29	Insufficient	Very low
Motor function of	Intra-observer reliability: Kappa: 0.50; total sample size: 29	Insufficient	Very low

shoulder and elbow	Inter-observer reliability: Kappa: 0.31; total sample size: 29		
Motor function of lower extremity	Intra-observer reliability: Kappa: 0.55; total sample size: 29 Inter-observer reliability: Kappa: 0.49; total sample size: 29	Insufficient	Very low
Sensory function of lower extremity	Intra-observer reliability: Kappa: 0.44; total sample size: 29 Inter-observer reliability: Kappa: 0.34; total sample size: 29	Insufficient	Very low
Sensory function of trunk	Intra-observer reliability: Kappa: 0.54; total sample size: 29 Inter-observer reliability: Kappa: 0.58; total sample size: 29	Insufficient	Very low
Sensory function of upper extremity	Intra-observer reliability: Kappa: 0.51; total sample size: 29 Inter-observer reliability: Kappa: 0.42; total sample size: 29	Insufficient	Very low
JOACMEQ			
Bladder function	Test-retest stability: ICC: 0.62; total sample size: 70	Insufficient	Very low
Cervical spine function	Test-retest stability: ICC: 0.63; total sample size: 70	Insufficient	Very low
Lower extremity function	Test-retest stability: ICC: 0.93; total sample size: 70	Sufficient	Very low
QOL	Test-retest stability:	Sufficient	Very low

		ICC: 0.83; total sample size: 70		
	Upper extremity function	Test-retest stability: ICC: 0.93; total sample size: 70	Sufficient	Very low
mJOA		Test-retest stability: Spearman's rank correlation: 0.91; total sample size: 75	Indeterminate	Very low
		Intra-observer reliability: ICC: 0.87; total sample size: 55	Sufficient	Very low
		Inter-observer reliability: ICC: 0.97; total sample size: 55 Kappa: 0.80; total sample size: 75	Sufficient	Low
	Motor dysfunction of lower extremities	Inter-observer reliability: ICC: 0.73; total sample size: 75	Sufficient	Low
	Motor dysfunction of upper extremities	Inter-observer reliability: ICC: 0.77; total sample size: 75	Sufficient	Low
	Sensory dysfunction of sphincter dysfunction	Inter-observer reliability: ICC: 0.78; total sample size: 75	Sufficient	Low
	Sensory dysfunction of upper extremities	Inter-observer reliability: ICC: 0.93; total sample size: 75	Sufficient	Low
MRI	(Depiction of intramedullary hyperintensity at eight cervical disc	Inter-observer reliability: Kendall's W range 0.72–0.78; total sample size: 79	Indeterminate	Very low

levels, T2W, 1.5-T or 3-T)			
MRI (Kang's classification, 1.5-T or 3-T)	Intra-observer reliability: Kappa: 0.67; total sample size: 439 ICC: 0.77, total sample size: 82 Inter-observer reliability: Kappa range 0.60–0.93; total sample size: 539 ICC range 0.74–0.75; total sample size: 82	Inconsistent	n/a
MRI (Muhle's classification, 1.5-T)	Intra-observer reliability: Kappa: 0.72; total sample size: 357 Inter-observer reliability: Kappa range 0.61; total sample size: 357	Inconsistent	n/a
MRI (Vaccaro's classification, 1.5-T)	Intra-observer reliability: Kappa: 0.71; total sample size: 357 Inter-observer reliability: Kappa range 0.69; total sample size: 357	Sufficient	Moderate
P-mJOA			
Motor dysfunction of lower extremities	Inter-observer reliability: Kappa: 0.61; total sample size: 755	Insufficient	Moderate
Motor dysfunction of upper extremities	Inter-observer reliability: Kappa: 0.66; total sample size: 755	Insufficient	Moderate
Sensory dysfunction of sphincter dysfunction	Inter-observer reliability: Kappa: 0.55; total sample size: 755	Insufficient	Moderate

Sensory dysfunction of upper extremities	Inter-observer reliability: Kappa: 0.55; total sample size: 755	Insufficient	Moderate
X-rays (Computer-assisted measurement of length)	Intra-observer reliability: ICC: 0.94; total sample size: 9 Inter-observer reliability: ICC: 0.93; total sample size: 9	Sufficient	Very low
X-rays (Computer-assisted measurement of thickness)	Intra-observer reliability: ICC: 0.96; total sample size: 9 Inter-observer reliability: ICC: 0.97; total sample size: 9	Sufficient	Very low

*Result ratings for BBS were split by statistic used due to their associated differences in sufficiency.

Supplementary Table 9. Measurement error.

Instrument	Result summary	Overall rating	Quality of evidence
BBS	MDC or SDC Distribution: 1.5; total sample size: 32	Indeterminate	n/a
EQ-5D	MDC or SDC Distribution: 0.13; total sample size: 101 Anchor: 0.04; total sample size: 101	Inconsistent	n/a
JOA	MDC or SDC Distribution: 1.0; total sample size: 101 Anchor: 2.5; total sample size: 101 LOA 1.2 (−1.2, 3.6); total sample size: 92	Sufficient	Very low
JOACMEQ			
Bladder function	MDC or SDC Distribution: 7.7; total sample size: 101	Insufficient	Very low
Cervical spine function	MDC or SDC Distribution: 12.9; total sample size: 101 Anchor: 12.5; total sample size: 101	Insufficient	Very low
Lower extremity function	MDC or SDC Distribution: 7.3; total sample size: 101 Anchor: 9.4; total sample size: 101	Inconsistent	n/a
QOL	MDC or SDC Distribution: 6.6; total sample size: 101 Anchor: 8.5; total sample size: 101	Sufficient	Very low

Upper extremity function	MDC or SDC Distribution: 9.5; total sample size: 101 Anchor: 6.1; total sample size: 101	Sufficient	Very low
mJOA	MDC or SDC Distribution: 2.1; total sample size: 113	Inconsistent	Very low
	MCID range; total sample size: 868 Distribution: 1.2–1.4	Sufficient	High
NDI	MDC or SDC Distribution: 6.2%; total sample size: 101 Anchor: 5.2%; total sample size: 101	Insufficient	Very low
SF-36			
MCS	MDC or SDC Distribution: 3.3–5.7; total sample size: 244	Inconsistent	n/a
	MCID; total sample size: 748 Distribution: 3.4–6.8	Inconsistent	n/a
PCS	MDC or SDC Distribution: 5.2–5.7; total sample size: 214 Anchor: 4.9; total sample size: 101	Inconsistent	n/a
	MCID range; total sample size: 861 Distribution: 2.9–5.5 MCID; total sample size: 51 Distribution: 10	Inconsistent	n/a
VAS for pain	MDC or SDC Distribution: 3.1; total sample size: 30	Insufficient	Very low

	MCID range 24.0–30.0; total sample size: 51	Insufficient	Very low
WHOQOL-Bref			
PH	MCID Distribution: 8.2; total sample size: 38	Indeterminate	n/a
PS	MCID Distribution: 7.9; total sample size: 38	Indeterminate	n/a
SR	MCID Distribution: 8.0; total sample size: 38	Indeterminate	n/a
EN	MCID Distribution: 5.6; total sample size: 38	Indeterminate	n/a
PF	MCID Distribution: 10.5; total sample size: 38	Indeterminate	n/a
RP	MCID Distribution: 17.2; total sample size: 38	Indeterminate	n/a
BP	MCID Distribution: 13.2; total sample size: 38	Indeterminate	n/a
GH	MCID Distribution: 12.3; total sample size: 38	Indeterminate	n/a
VT	MCID Distribution: 10.8; total sample size: 38	Indeterminate	n/a
SF	MCID Distribution: 13.6; total sample size: 38	Indeterminate	n/a
RE	MCID Distribution: 18.0; total sample size: 38	Indeterminate	n/a
MH	MCID	Indeterminate	n/a

Distribution: 11.2; total
sample size: 38

n/a = No info available

Supplementary Table 10. Criterion validity.

Instrument	Result summary*	Overall rating	Quality of evidence
10-s step test	JOA Spearman's rank correlation: 0.66; total sample size: 163	Insufficient	High
BBS	mJOA AUC range 0.88–0.94; total sample size: 31	Sufficient	Low
Foot tapping test	JOA Pearson's correlation: 0.66; total sample size: 126 JOA MFLE Pearson's correlation: 0.70; total sample size: 126	Insufficient	High
Grip-and-release test	JOA Pearson's correlation: 0.72; total sample size: 30	Sufficient	Low
JOA	Likert scale, "Health transition question" AUC: 0.59; total sample size: 101 Likert scale, "Patient satisfaction question" AUC: 0.62; total sample size: 101	Insufficient	Very low
JOACMEQ			
Cervical spine function	Likert scale, "Health transition question" AUC: 0.58; total sample size: 101 Likert scale, "Patient satisfaction question"	Insufficient	Very low

		AUC: 0.58; total sample size: 101		
Upper extremity function	Likert scale, "Health transition question" AUC: 0.66; total sample size: 101 Likert scale, "Patient satisfaction question" AUC: 0.65; total sample size: 101	Insufficient	Very low	
Lower extremity function	Likert scale, "Health transition question" AUC: 0.61; total sample size: 101 Likert scale, "Patient satisfaction question" AUC: 0.66; total sample size: 101	Insufficient	Very low	
QOL	Likert scale, "Health transition question" AUC: 0.70; total sample size: 101 Likert scale, "Patient satisfaction question" AUC: 0.66; total sample size: 101	Insufficient	Very low	
mJOA	Nurick scale [convergent] Spearman's rank correlation: -0.41; total sample size: 119 Pearson's correlation range: -0.62 to -0.63; total sample size: 352	Sufficient	High	
Motor dysfunction of upper extremities	Nurick scale [convergent] Pearson's correlation range -0.42 to -0.42; total sample size: 352	Insufficient	High	

Motor dysfunction of lower extremities	Nurick scale [convergent] Pearson's correlation: -0.65 to -0.68; total sample size: 352	Sufficient	High
Sensory dysfunction of upper extremities	Nurick scale [convergent] Pearson's correlation: -0.23; total sample size: 277	Insufficient	High
Sensory dysfunction of sphincter dysfunction	Nurick scale [convergent] Pearson's correlation: -0.25; total sample size: 277	Insufficient	High
NDI	Likert scale, "Health transition question" AUC: 0.66; total sample size: 101 Likert scale, "Patient satisfaction question" AUC: 0.75; total sample size: 101	Inconsistent	n/a
P-mJOA	mJOA Spearman's rank correlation: 0.83; total sample size: 755	Sufficient	High
QuickDASH	JOA MFSE Spearman's rank correlation: -0.50; total sample size: 94 JOA SFUE Spearman's rank correlation: -0.32; total sample size: 94	Insufficient	Moderate
SF-36			
PCS	Likert scale, "Health transition question" AUC: 0.67; total sample size: 101	Insufficient	Very low

		Likert scale, "Patient satisfaction question" AUC: 0.69; total sample size: 101		
WHOQOL-Bref				
PH	SF-36 PCS	Pearson's correlation: 0.51; total sample size: 38	Inconsistent	n/a
	SF-36 MCS	Pearson's correlation: 0.30; total sample size: 38		
PS	SF-36 PCS	Pearson's correlation: 0.34; total sample size: 38	Insufficient	Low
	SF-36 MCS	Pearson's correlation: 0.23; total sample size: 38		
SR	SF-36 PCS	Pearson's correlation: 0.35; total sample size: 38	Insufficient	Low
	SF-36 MCS	Pearson's correlation: 0.28; total sample size: 38		
EN	SF-36 PCS	Pearson's correlation: 0.05; total sample size: 38	Insufficient	Low
	SF-36 MCS	Pearson's correlation: 0.03; total sample size: 38		

n/a = No info available

*Instruments listed are comparators

Supplementary Table 11. Construct validity.

Instrument	Result summary*	Overall rating	Quality of evidence
10-s step test	Grip-and-release test [convergent] Spearman's rank correlation: 0.53; total sample size: 163	Sufficient	Moderate
30MWT	mJOA [convergent] Pearson's correlation: -0.44; total sample size: 16 MDI [convergent] Spearman's rank correlation: 0.65; total sample size: 41 Nurick scale [convergent] Pearson's correlation: 0.50; total sample size: 16 Spearman's rank correlation: 0.61; total sample size: 41	Sufficient	Moderate
	NDI Pearson's correlation: 0.21; total sample size: 16	Sufficient	Low
	SF-36 PCS Pearson's correlation: -0.35; total sample size: 16	Sufficient	Low
	SF-36 MCS Pearson's correlation: -0.20; total sample size: 16	Sufficient	Low
BBS	mJOA [convergent] Spearman's rank correlation: 0.81; total sample size: 72	Sufficient	Moderate

EQ-5D	mJOA AUC: 0.68; total sample size: 119 Nurick scale AUC: 0.61; total sample size: 119	Insufficient	High
Foot tapping test	Grip-and-release test [convergent] Pearson's correlation: 0.58; total sample size: 126	Sufficient	High
Isihara's Cervical Curvature Index	mJOA Pearson's correlation: 0.04; total sample size: 235	Sufficient	High
	SF-12 PCS Pearson's correlation: 0.06; total sample size: 235	Sufficient	High
	SF-12 MCS Pearson's correlation: 0.11; total sample size: 235		
	Pain, "Numeric rating scale" (Arm pain score) Pearson's correlation: -0.28; total sample size: 235	Sufficient	High
	Pain, "Numeric rating scale" (Neck pain scores) Pearson's correlation: -0.27; total sample size: 235		
	NDI Pearson's correlation: -0.10; total sample size: 235		
JOA	mJOA [convergent]	Sufficient	Low

		Spearman's rank correlation: 0.87; total sample size: 92		
		JOACMEQ QOL [convergent]	Sufficient	Low
		Spearman's rank correlation: 0.41; total sample size: 92		
		SF-12 PCS	Sufficient	Low
		Spearman's rank correlation: 0.50; total sample size: 92		
		SF-12 MCS	Sufficient	Low
		Spearman's rank correlation: -0.05; total sample size: 92		
		NDI	Sufficient	Moderate
		Spearman's rank correlation range – 0.50 to -0.76; total sample size: 122		
JOACMEQ				
	QOL	NDI	Sufficient	Low
		Spearman's rank correlation: -0.66; total sample size: 92		
	QOL	SF-12 PCS	Insufficient	Low
		Spearman's rank correlation: 0.29; total sample size: 92		
		SF-12 MCS		
		Spearman's rank correlation: 0.40; total sample size: 92		
	MDI	Nurick scale [convergent]	Sufficient	Low
		Spearman's rank correlation: 0.66; total sample size: 41		

mJOA	30MWT [convergent]	Insufficient	High
	Pearson's correlation: –0.38; total sample size: 193		
	JOACMEQ QOL [convergent]	Insufficient	Low
	Spearman's rank correlation: 0.41; total sample size: 92		
	EQ-5D	Insufficient	High
	Spearman's rank correlation: 0.42; total sample size: 119		
	SF-36 PCS	Sufficient	High
Pearson's correlation range: 0.30–0.30; total sample size: 338			
SF-12 PCS			
Spearman's rank correlation: 0.47; total sample size: 92			
SF-36 MCS			
Sufficient			
High			
Pearson's correlation: 0.25–0.25; total sample size: 338			
SF-12 MCS			
Spearman's rank correlation: 0.03; total sample size: 92			
NDI			
Sufficient			
High			
Spearman's rank correlation: –0.51; total sample size: 92			
Pearson's correlation range –0.33 to –0.34; total sample size: 336			
Motor dysfunction	30MWT [convergent]	Insufficient	High

of lower extremities	Pearson's correlation: –0.43; total sample size: 193		
	SF-36 PCS	Sufficient	High
	Pearson's correlation range: 0.31–0.50; total sample size: 338		
	SF-36 MCS	Sufficient	High
	Pearson's correlation: 0.21; total sample size: 268		
	NDI	Sufficient	High
	Pearson's correlation: –0.31; total sample size: 261		
	Motor dysfunction of upper extremities	30MWT [convergent]	Insufficient
	Pearson's correlation: –0.21; total sample size: 193		
	SF-36 PCS	Insufficient	High
	Pearson's correlation: 0.22; total sample size: 268		
	SF-36 MCS	Sufficient	High
	Pearson's correlation: 0.20; total sample size: 268		
	NDI	Sufficient	High
	Pearson's correlation: –0.24; total sample size: 261		
	Sensory dysfunction of sphincter dysfunction	30MWT [convergent]	Insufficient
	Pearson's correlation: –0.23; total sample size: 193		
	SF-36 PCS	Sufficient	High

		Pearson's correlation: 0.06; total sample size: 268		
		SF-36 MCS Pearson's correlation: 0.08; total sample size: 268		
	NDI		Sufficient	High
		Pearson's correlation: -0.16; total sample size: 261		
Sensory dysfunction of upper extremities	30MWT [convergent]		Insufficient	High
		Pearson's correlation: -0.05; total sample size: 193		
	SF-36 PCS		Sufficient	High
		Pearson's correlation: 0.19; total sample size: 268		
	SF-36 MCS			
		Pearson's correlation: 0.19; total sample size: 268		
	NDI		Insufficient	High
		Pearson's correlation: -0.23; total sample size: 261		
NDI	mJOA		Sufficient	High
		Pearson's correlation: -0.36; total sample size: 235		
	SF-12 PCS		Sufficient	High
		Pearson's correlation: -0.54; total sample size: 235		
	SF-12 MCS			

		Pearson's correlation: -0.40; total sample size: 235		
	Pain, "Numeric rating scale" (Arm pain score)		Sufficient	High
		Pearson's correlation: 0.68; total sample size: 235		
	Pain, "Numeric rating scale" (Neck pain scores)			
		Pearson's correlation: 0.64; total sample size: 235		
Nurick scale	EQ-5D		Sufficient	High
		Spearman's rank correlation: -0.28; total sample size: 119		
Pain, "Numeric rating scale" (Arm pain scores)	mJOA		Sufficient	High
		Pearson's correlation: -0.19; total sample size: 235		
	Pain, "Numeric rating scale" (Neck pain score) [convergent]		Sufficient	High
		Pearson's correlation: 0.72; total sample size: 235		
Pain, "Numeric rating scale" (Neck pain scores)	mJOA		Sufficient	High
		Pearson's correlation: -0.07; total sample size: 235		
QuickDASH	SF-36		Sufficient	Moderate
		Spearman's rank correlation: -0.75; total sample size: 94		
	NDI and Pain, "Numeric rating scale" [convergent]		Sufficient	Moderate

		Spearman's rank correlation range 0.69–0.83; total sample size: 94		
SF-12				
MCS	mJOA		Sufficient	High
		Pearson's correlation: 0.19; total sample size: 235		
	Pain, "Numeric rating scale" (Arm pain score)		Sufficient	High
		Pearson's correlation: –0.23; total sample size: 235		
	Pain, "Numeric rating scale" (Neck pain score)			
		Pearson's correlation: –0.28; total sample size: 235		
	NDI		Sufficient	Moderate
		Spearman's rank correlation: –0.17; total sample size: 92		
	SF-12 PCS		Sufficient	High
		Pearson's correlation: 0.01; total sample size: 235		
PCS	mJOA		Sufficient	High
		Pearson's correlation: 0.43; total sample size: 235		
	Pain, "Numeric rating scale" (Arm pain score)		Sufficient	High
		Pearson's correlation: –0.44; total sample size: 235		
	Pain, "Numeric rating scale" (Neck pain score)			

	Pearson's correlation: -0.41; total sample size: 235		
	NDI	Sufficient	Moderate
	Spearman's rank correlation: -0.49; total sample size: 92		
	SF-12 MCS	Sufficient	Low
	Spearman's rank correlation: -0.29; total sample size: 92		
Triangle step test	Grip-and-release test [convergent]	Sufficient	High
	Spearman's rank correlation: 0.55; total sample size: 270		

*Instruments listed are comparators

Supplementary Table 12. Responsiveness.

Instrument	Result summary*	Overall rating	Quality of evidence
30MWT	30MWT SRM: 0.3; total sample size: 484	Insufficient	High
BBS	mJOA Sensitivity range 77.4-80.0; total sample size: 31 Specificity range 87.8-92.9; total sample size: 31	Sufficient	Low
EQ-5D	EQ-5D Mean change score: 0.06; total sample size: 108	Indeterminate	High
European Myelopathy Scale	EMS Normalised change: 0.18; total sample size: 99	Indeterminate	Very low
Foot tapping test	Foot tapping test Mean change score: 6; total sample size: 6	Indeterminate	Very low
Grip-and-release test	Grip-and-release test Spearman's rank correlation: 0.69; total sample size: 48	Sufficient	Very low
	JOA Spearman's rank correlation: 0.32; total sample size: 48	Insufficient	Low
JOA	mJOA Spearman's rank correlation: 0.75; total sample size: 92	Sufficient	Very low
	JOA	Indeterminate	Very low

		Mean change score range 4.6; total sample size: 126 Normalised change: 0.21; total sample size: 99		
		JOA MFLE Mean change score range 0.6; total sample size: 126		
JOACMEQ				
Bladder function	JOACMEQ BF		Sufficient	Moderate
		AUC: 0.82; total sample size: 78		
	JOACMEQ BF		Indeterminate	Very low
		Mean change score: 18.0; total sample size: 87		
	JOACMEQ BF		Insufficient	Moderate
		Effect size: 0.33; total sample size: 78		
Cervical spine function	JOACMEQ CF		Sufficient	Moderate
		AUC: 0.72; total sample size: 78		
	JOACMEQ CF		Indeterminate	Very low
		Mean change score: 25.8; total sample size: 87		
	JOACMEQ CF		Insufficient	Moderate
		Effect size: 0.28; total sample size: 78		
Lower extremity function	JOACMEQ LEF		Sufficient	Moderate
		AUC: 0.75; total sample size: 78		
	JOACMEQ LEF		Indeterminate	Very low
		Mean change score: 28.4; total sample size: 87		

	JOACMEQ LEF	Insufficient	Moderate
	Effect size: 0.02; total sample size: 78		
Upper extremity function	JOACMEQ UEF	Sufficient	Moderate
	AUC: 0.74; total sample size: 78		
	JOACMEQ UEF	Indeterminate	Very low
	Mean change score: 10.7; total sample size: 87		
	JOACMEQ UEF	Insufficient	Moderate
	Effect size: 0.17; total sample size: 78		
QOL	JOACMEQ QOL	Sufficient	Moderate
	AUC: 0.83; total sample size: 78		
	JOACMEQ QOL	Indeterminate	Very low
	Mean change score: 23.7; total sample size: 87		
	JOACMEQ QOL	Insufficient	Moderate
	Effect size: 0.46; total sample size: 78		
MDI	MDI	Indeterminate	Very low
	Normalised change: 0.52; total sample size: 99		
mJOA	mJOA	Sufficient	High
	Effect size range 0.87–1.0; total sample size: 352		
	mJOA	Indeterminate	Very low
	Normalised change: 1.47; total sample size: 42		
NDI	Anchor-based approach	Insufficient	Moderate
	AUC: 0.66; total sample size: 78		

		Effect size: 0.44; total sample size: 78		
	NDI		Indeterminate	Very low
		Mean change score: –15.8; total sample size: 118		
Nurick scale	Nurick scale		Indeterminate	Very low
		Normalised change: 0.42; total sample size: 99		
		Mean change score range –0.76 to –1.3; total sample size: 93		
Ranawat classification of disease severity	Ranawat classification of disease severity		Indeterminate	Very low
		Normalised change: 0.34; total sample size: 99		
SF-12				
	PCS	SF-12 PCS	Indeterminate	Very low
		Mean change score: 8.17; total sample size: 118		
SF-36	SF-36		Indeterminate	Very low
		Normalised change: 0.32; total sample size: 99		
	PCS	SF-36 PCS	Sufficient	Low
		Effect size range: 0.84; total sample size: 142		
		SF-36 PCS	Sufficient	Moderate
		Sensitivity: 0.85; total sample size: 105		
	MCS	SF-36 MCS	Sufficient	Low
		Effect size range: 0.81; total sample size: 142		
		SF-36 MCS	Sufficient	Moderate

Sensitivity: 0.67; total sample size: 105			
WHOQOL-Bref			
PH	WHOQOL-Bref PH	Insufficient	Low
	Effect size: 0.68; total sample size: 38		
PS	WHOQOL-Bref PS	Insufficient	Low
	Effect size: 0.39; total sample size: 38		
SR	WHOQOL-Bref SR	Insufficient	Low
	Effect size: 0.03; total sample size: 38		
EN	WHOQOL-Bref EN	Insufficient	Low
	Effect size: 0.45; total sample size: 38		

*Instruments listed are comparators

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