

University of Birmingham Research at Birmingham

Physical functioning in adolescents with idiopathic scoliosis

Alamrani, Samia; Rushton, Ali; Gardner, Adrian; Bini, Elena; Falla, Deborah; Heneghan, Nicola

DOI:

10.1097/BRS.0000000000003969

License:

Creative Commons: Attribution-NonCommercial (CC BY-NC)

Document Version
Peer reviewed version

Citation for published version (Harvard):

Alamrani, S, Rushton, A, Gardner, A, Bini, E, Falla, D & Heneghan, N 2021, 'Physical functioning in adolescents with idiopathic scoliosis: a systematic review of outcome measures and their measurement properties', *Spine*, vol. 46, no. 18, pp. E985-E997. https://doi.org/10.1097/BRS.000000000003969

Link to publication on Research at Birmingham portal

General rights

Unless a licence is specified above, all rights (including copyright and moral rights) in this document are retained by the authors and/or the copyright holders. The express permission of the copyright holder must be obtained for any use of this material other than for purposes permitted by law.

•Users may freely distribute the URL that is used to identify this publication.

•Users may download and/or print one copy of the publication from the University of Birmingham research portal for the purpose of private study or non-commercial research.

•User may use extracts from the document in line with the concept of 'fair dealing' under the Copyright, Designs and Patents Act 1988 (?)

•Users may not further distribute the material nor use it for the purposes of commercial gain.

Where a licence is displayed above, please note the terms and conditions of the licence govern your use of this document.

When citing, please reference the published version.

Take down policy

While the University of Birmingham exercises care and attention in making items available there are rare occasions when an item has been uploaded in error or has been deemed to be commercially or otherwise sensitive.

If you believe that this is the case for this document, please contact UBIRA@lists.bham.ac.uk providing details and we will remove access to the work immediately and investigate.

Download date: 23. Apr. 2024

Spine (Phila Pa 1976)

Physical functioning in adolescents with idiopathic scoliosis: A systematic review of outcome measures and their measurement properties --Manuscript Draft--

Manuscript Number:	SPINE 162799
Full Title:	Physical functioning in adolescents with idiopathic scoliosis: A systematic review of outcome measures and their measurement properties
Article Type:	Literature Review
Keywords:	Systematic Review; Idiopathic Scoliosis; Physical Functioning; Outcome Assessment; Validity; Reliability; Measurement Properties.
Corresponding Author:	Samia Alamrani, MSc University of Birmingham Birmingham, West Midland UNITED KINGDOM
Corresponding Author Secondary Information:	
Corresponding Author's Institution:	University of Birmingham
Corresponding Author's Secondary Institution:	
First Author:	Samia Alamrani, MSc
First Author Secondary Information:	
Order of Authors:	Samia Alamrani, MSc
	Alison B Rushton, EdD
	Elena Bini, MSc
	Adrian Gardner, PhD
	Deborah Falla, PhD
	Nicola R Heneghan, PhD
Order of Authors Secondary Information:	
Additional Information:	
Question	Response
Please provide the Word Count of your manuscript text. Include only the main body of text (exclude abstract, references, figures, and table legends).	2686
Please provide the Word Count of your structured abstract. Include only the abstract, not key words.	299
Please select the <u>level of evidence</u> for this manuscript.	2
Device Status/Drug Statement: Please select the statement below that applies to your submission. After you have selected the appropriate statement, Please also add the same statement to the Title Page of your submission.	The Manuscript submitted does not contain information about medical device(s)/drug(s).

RETAINED RIGHTS: Except for copyright, other proprietary rights related to

the Work (e.g., patent or other rights to any process or procedure) shall be retained by the author. To reproduce any text, figures, tables, or illustrations from this Work in future works of their own, the author must obtain written permission from Wolters Kluwer Health, Inc. ("WKH").

ORIGINALITY: Each author warrants that his or her submission to the Work is original, does not infringe upon, violate, or misappropriate any copyright or other intellectual property rights, or any other proprietary right, contract or other right or interest of any third party, and that he or she has full power to enter into this agreement. Neither this Work nor a similar work has been published nor shall be submitted for publication elsewhere while under consideration by this Publication.

AUTHORSHIP RESPONSIBILITY: Each author warrants that he or she has participated sufficiently in the intellectual content, the analysis of data, if applicable, and the writing of the Work to take public responsibility for it. Each has reviewed the final version of the Work, believes it represents valid work, and approves it for publication. Moreover, should the editors of the Publication request the data upon which the work is based, they shall produce it.

PREPRINTS: Upon acceptance of the article for publication, each author warrants that he/she will promptly remove any prior versions of this Work (normally a preprint) that may have been posted to an electronic server.

I agree

DISCLAIMER: Each author warrants that this Work contains no libelous or unlawful statements and does not infringe or violate the publicity or privacy rights of any third party, libel or slander any third party, contain any scandalous, obscene, or negligently prepared information, or infringe or violate any other personal or proprietary right of others. Each author warrants that the Work does not contain any fraudulent, plagiarized or incorrectly attributed material. Each author warrants that all statements contained in the Work purporting to be facts are true, and any formula or instruction contained in the Work will not, if followed accurately, cause any injury, illness, or damage to the user. If excerpts (e.g., text, figures, tables, illustrations, or audio/video files) from copyrighted works are included, a written release will be secured by the author prior to submission, and credit to the original publication will be properly acknowledged. Each author further warrants that he or she has obtained, prior to submission, written releases from patients whose names or likenesses are submitted as part of the Work. Should the Editor or WKH request copies of such written releases, the author shall provide them in a timely manner.

DISCLOSURES/CONFLICT OF INTEREST

Each author must identify any financial interests or affiliations with institutions, organizations, or companies relevant to the manuscript by completing the form below. Additionally, any financial associations involving a spouse, partner or children must be disclosed as well.

Note: Some sections below come from the ICMJE Uniform Disclosure Form for Potential Conflicts of

Interest at http://www.icmje.org/downloads/coi_disclo	
sure.pdf (dated July 2010).	
Did you or your institution at any time receive payment or support in kind for any aspect of the submitted work (including but not limited to grants, consulting fee or honorarium, support for travel to meetings for the study or other purposes, fees for participation in review activities such as data monitoring boards, statistical analysis, end point committees, and the like, payment for writing or reviewing the manuscript, provision of writing assistance, medicines, equipment, or administrative support, etc)?	No
Other: Did you or your institution at any time receive additional payments or support in kind for any aspect of the submitted work?	None
Please indicate whether you have financial relationships (regardless of amount of compensation) with entities. You should report relationships that were present during the 36 months prior to submission including board membership, consultancy, employment, expert testimony, grants/grants pending, payment for lectures including service on speakers bureaus, payment for manuscript preparation, patents (planned, pending or issued), royalties, payment for development of educational presentations, stock/stock options, travel/accommodations/meeting expenses unrelated to activities listed (for example, if you report a consultancy above there is no need to report travel related to that consultancy), etc.	No No
Other (err on the side of full disclosure): Please indicate whether you have any additional financial relationships (regardless of amount of compensation) with entities. You should report relationships that were present during the 36 months prior to submission.	
Other Relationships Are there other relationships or activities that readers could perceive to have	No other relationships/conditions/circumstances that present potential conflict of interest

influenced, or that give the appearance of potentially influencing, what you wrote in the submitted work?

AUTHOR'S OWN WORK: In

consideration of WKH's publication of the Work, the author hereby transfers, assigns, and otherwise conveys all his/her copyright ownership worldwide, in all languages, and in all forms of media now or hereafter known, including electronic media such as CD-ROM, Internet, and Intranet, to WKH. If WKH should decide for any reason not to publish the Work, WKH shall give prompt notice of its decision to the corresponding author, this agreement shall terminate, and neither the author nor WKH shall be under any further liability or obligation. Each author grants WKH the rights to use his or her name and biographical data (including professional affiliation) in the Work and in its or the journal's promotion. Notwithstanding the foregoing, this paragraph shall not apply, and any transfer made pursuant to this paragraph shall be null and void if (i) the Work has been accepted by WKH for publication, and (ii) the author chooses to have the Work published by WKH as an open access publication.

WORK MADE FOR HIRE: If this Work or any element thereof has been commissioned by another person or organization, or if it has been written as part of the duties of an employee, an authorized representative of the commissioning organization or employer must also sign this form stating his or her title in the organization.

GOVERNMENT EMPLOYEES: If the Work or a portion of it has been created in the course of any author's employment by the United States Government, check the "Government" box at the end of this form. A work prepared by a government employee as part of his or her official duties is called a "work of the U.S. Government" and is not subject

I agree

to copyright. If it is not prepared as part of the employee's official duties, it may be subject to copyright. **INSTITUTIONAL REVIEW BOARD/ANIMAL CARE COMMITTEE** APPROVAL: Each author warrants that his or her institution has approved the protocol for any investigation involving humans or animals and that all experimentation was conducted in conformity with ethical and humane principles of research. WARRANTIES: Each author warranty made in this form is for the benefit of WKH and the Editor; each author agrees to defend, indemnify, and hold harmless those parties for any breach of such warranties. Spine will permit the author(s) to deposit I agree for display a "final peer-reviewed manuscript" (the final manuscript after peer-review and acceptance for publication but prior to the publisher's copyediting, design, formatting, and other services) 12 months after publication of the final article on the author's personal web site, university's institutional repository or employer's intranet, subject to the following: * You may only deposit the final peerreviewed manuscript. * You may not update the final peerreviewed manuscript text or replace it with proof or with the final published version. * You may not include the final peerreviewed manuscript or any other version the article on any commercial site or in any repository owned or operated by any third party. For authors of articles

based on research funded by the National

Institutes of Health ("NIH"), Wellcome Trust, Howard Hughes Medical Institute ("HHMI"), or other funding agency, see below for the services that WKH will provide on your behalf to comply with "Public Access Policy" guidelines.	
* You may not display the final peer- reviewed manuscript until twelve months after publication of the final article.	
* You must attach the following notice to the final peer-reviewed manuscript: "This is a non-final version of an article published in final form in (provide complete journal citation)".	
* You shall provide a link in the final peer-reviewed manuscript to the Spine website.	
"Public Access Policy" Funding Disclosure Please disclose below if you have received funding for research on which your article is based from any of the following organizations:	
Please select:	Author's Own Work
Any additional comments?	
Compliance with RCUK and Wellcome Trust Open Access Policies Both the Research Councils UK (RCUK)	I agree
and the Wellcome Trust have adopted policies regarding Open Access to articles that have been funded by grants from the RCUK or the Wellcome Trust. If either "Wellcome Trust" or "Research Councils UK (RCUK)" has been selected above, and	
the authors of the applicable article choose to have the article published as an open access publication, the following policies will apply:	

Wellcome Trust require that WKH make the article freely available immediately pursuant to the Attribution 4.0 Creative Commons License, currently found at

http://creativecommons.org/licenses/by/4. 0/legalcode

(the "CC BY License"). The CC BY License is the most accommodating of the Creative Commons licenses and allows others to distribute, remix, tweak, and build upon the article, even commercially, as long as they credit the authors for the original creation.

* If the article is to be published pursuant to the "Green" route of Open Access, both the RCUK and the Wellcome Trust require that WKH make the article freely available within six months pursuant to the Attribution-NonCommerical 4.0 Creative Commons License, currently found at http://creativecommons.org/licenses/bync/4.0/legalcode (the "CC BY-NC License"). The CC BY-NC License allows others to remix, tweak, and build upon the article noncommercially, and although their new works must also acknowledge the authors for the original creation and be non-commercial, they don't have to license their derivative works on the same terms.

As a service to our authors, WKH will identify the National Library of Medicine (NLM) articles that require deposit pursuant to the RCUK and Wellcome Trust policies described in this section. This Copyright Transfer Agreement provides the mechanism for identifying such articles.

WKH will transmit the final peer-reviewed manuscript of an article based on research funded in whole or in part by either RCUK or the Wellcome Trust to

Pub Med Central. Upon NIH request, it remains the legal responsibility of the author to confirm with NIH the provenance of his/her manuscript for purposes of deposit. Author will not deposit articles him/herself. Author will not alter the final peer-reviewed manuscript already transmitted to NIH. With respect to the "Green" route of Open Access, author will not authorize the display of the final peer-reviewed manuscript prior to 6 months following publication of the final article. Authors of articles that have been funded from grants from the RCUK or the Wellcome Trust are required to sign the WKH Open Access License Agreement prior to publication of the applicable article. Please contact the Editorial Office of the applicable journal to receive the Open Access License Agreement that is to be signed in connection with the publication of the article. I am the person in question for this I agree submission or otherwise have approval to complete this agreement. CME/CE Disclosure I agree Each author must identify and disclose any financial associations involving a spouse, partner or children by completing the Family Disclosure question below, and whether any off-label uses or unapproved drugs or devices are discussed in his/her manuscript by completing the Off-Label Use/Unapproved Drugs or Products question below. In the event that the Work is published as a continuing education or continuing medical education article, this information will be provided to the accrediting body and may be included in the published

article. When applicable, articles accepted for publication may need to comply with additional standards related to CME or CE accreditation. Please refer to guidelines for authors for details. WKH and its affiliates reserve the right to publish the manuscript as a continuing education article.	
Pamily Disclosure Do your children or your spouse or partner have financial relationships with entities that have an interest in the content of the submitted work?	No other relationships/conditions/circumstances that present potential conflict of interest
Off-Label Use/Unapproved Drugs or Products If your manuscript discusses an unlabeled use of a commercial product or device or an investigational use of a product or device not yet approved by the FDA for any purpose, you must specifically disclose in the manuscript that the product is not labeled for the use under discussion or that the product is still investigational. Please check the item below that applies to you	I will not discuss unlabeled/investigational uses of any commercial product or device

Physical functioning in adolescents with idiopathic scoliosis: A systematic review of outcome measures and their measurement properties.

Samia Alamrani (MSc)^{1, 2}, Alison B Rushton (EdD) ^{1,3}, Elena Bini (MSc)¹, Adrian Gardner (PhD) ⁴, Deborah Falla (PhD) ¹, Nicola R Heneghan (PhD) ¹

¹ Centre of Precision Rehabilitation for Spinal Pain (CPR Spine), School of Sport, Exercise and Rehabilitation Sciences, College of Life and Environmental Sciences, University of Birmingham, Birmingham, United Kingdom

² Physical Therapy Department, College of Applied Medical Science, University of Tabuk, Tabuk, Saudi Arabia

³ School of Physical Therapy, Western University, London, Ontario, Canada

⁴ Spine Unit, The Royal Orthopaedic Hospital, Northfield, Birmingham, United Kingdom

Address correspondence and reprint requests to Samia Alamrani, PT, MSc, PhD student at Centre of Precision Rehabilitation for Spinal Pain (CPR Spine), School of Sport, Exercise and Rehabilitation Sciences, College of Life and Environmental Sciences, University of Birmingham, Birmingham, United Kingdom. E-mail: sxa1258@student.bham.ac.uk; salamrani@ut.edu.sa

The manuscript submitted does not contain information about medical device(s)/drug(s).

No funds were received in support of this work. SA is a PhD student had a scholarship from University of Tabuk, Tabuk, Saudi Arabia

No relevant financial activities outside the submitted work.

Abstract

Study Design. A systematic review

Objective. To summarise evidence on measurement properties of outcome measures (OM) used to assess physical functioning in Adolescents with Idiopathic Scoliosis (AIS).

Summary of Background Data. The AIS is a common spine deformity in those aged 10 to 18 years old. Associated health problems (e.g., back pain) significantly impact the quality of life (QoL). One important domain in QoL is physical functioning, which can be measured with Patient-Reported Outcome Measures (PROM), Performance-Based Outcome Measures (PBOM), and body structure and function OM. Adequate measurement properties of outcome measures (OM) are important for precision in research and practice

Methods. A two-search strategy performed on electronic databases up to December 2019. Search one revealed list of OM were used for physical functioning assessment in AIS. Search two identified studies that evaluated measurement property in AIS; using list identified in search one. Two independent reviewers determined study eligibility, risk of bias assessment (COSMIN checklist), and data extraction. The level of evidence was established using modified GRADE approach.

Results: Search one yielded: 28 PROM, 20 PBOM, and 10 body structure and function OM. Search two revealed: 16 measurement properties studies of PROM, 1 for PBOM and 3 for body structure and function measure. Construct validity, reliability and responsiveness of most PROMs established in AIS, but not content validity or internal consistency (Moderate evidence). Construct validity was sufficient for the Timed up and Go test and, body structure and function measures (very low to low evidence).

Conclusion: Currently, physical functioning evaluated with variety of measures in AIS. Majority of measurement properties studies were evaluating PROM with paucity of information on measurement properties of PBOM and body structure and function OM. Based

on COSMIN methodology, none of OM identified in this review can be recommended for use in individuals with AIS.

Key Words: Systematic Review, Idiopathic Scoliosis, Physical Functioning, Outcome Assessment, Validity, Reliability, Measurement Properties.

Level of Evidence: 2

Key points

- Two searched-strategy performed on all types of outcome measure used in physical functioning assessment for Adolescent with Idiopathic Scoliosis (AIS).
- Most of studies of measurement properties were evaluating Patient Reported Outcome Measure (PROM) with paucity of information on Performance-Based Outcome Measure (PBOM), and body structure and function measures.
- Based on COSMIN methodology, none of measure identified in this review can be recommended for use in individuals with AIS.

Mini Abstract (50 words)

Mini abstract

This review identified a variety of outcome measure used for physical functioning assessment in Adolescent with Idiopathic Scoliosis. However, a limited number of studies evaluated its measurement properties with focus was on patient-reported outcome measure compared to other outcome measures i.e. performance-based and body structure and function measures.

INTRODUCTION

Adolescent idiopathic scoliosis (AIS) is the most common spine deformity among children aged 10 to 18 years old ¹, with prevalence ranging 1-3% ². Comprising of a lateral curvature and axial rotation of spinal vertebrae, the cause is unknown in most cases ³. AIS has been linked to back pain ⁴, psychological stress ⁵, and respiratory dysfunction ⁶, potentially impacting on quality of life (QoL) ⁷. A dimension of any QoL measurement is 'physical functioning', this being the ability to carry out activities of daily living 8. Physical functioning limitations have been associated with an increased risk of disability and predictive of social and healthcare use 9. Limitations include walking and maintaining body positions ⁷, as well as pain related functional restriction ¹⁰. Corrective surgery is used for some, necessitating a long recovery period and often associated with pain and immobility in adolescence ¹¹. Measuring the impact of AIS is therefore important in both research and clinical practice. Physical functioning can be evaluated with Patient-Reported Outcome Measures (PROM), Performance-Based Outcome Measures (PBOM), and measures of body structure and function ¹². Each measure assesses different, but complementary, aspects of physical functioning ¹², with PROM for self-report, PBOM for the performance of a specific activity (e.g., chair stand test) ^{12,13} and body structure and function providing anatomical data (e.g., range of motion) or a physiological process (e.g., muscle strength) ¹². Outcome measures need adequate measurement properties to assure truthfulness of results and avoid risk of bias ¹⁴. The COnsensus-based Standards for the selection of health Measurement INstruments (COSMIN) group developed a taxonomy of measurement properties to enable this ¹⁵. Three main domains are validity, reliability and responsiveness ¹⁵. The COSMIN group

 provide guidelines for conducting a systematic review for PROM, which can be adapted for other OM ¹⁶.

The Scoliosis Research Society questionnaire (SRS-22) and its' variants are the most widely used PROM in this population ¹⁷⁻¹⁹. From the Core Outcome Study (COS), SRS-22 revised (SRS-22r) is recommended and the considered reference standard for evaluating physical functioning for adolescents and young adults with spine deformity ²⁰. However, SRS-22r does not capture all aspects of physical functioning, such as mobility and self-care ⁷. Furthermore, the COS study included all forms of spinal deformities; the heterogeneity limiting applicability to individuals with AIS. Furthermore, little is known about PBOM and body structure and

In the absence of existing relevant reviews, ²¹. the purpose of this review was to identify OM used to assess physical functioning in individuals with AIS, and secondly to evaluate their measurement properties.

Methods

function measures for individuals with AIS.

Design

This review was conducted according to a registered (PROSPERO CRD42019142335) and published protocol ²². Designed in line with COSMIN methodology for systematic review of PROM ¹⁶, the review is reported in line with Preferred Reporting Items for Systematic Review and Meta-Analysis (PRISMA) statement ²³.

Search strategy

The search was conducted in two parts. Search one identified and generated a list of OM used for assessment of physical functioning in AIS. Search two identified the studies of measurement properties using the list from search one. Details of both search are listed in Table 1.

Data sources

 A comprehensive search was performed using MEDLINE, PsycINFO, EMBASE, CINAHL, SPORTdiscus, Web of Science and PubMed databases from date of inception until December 2019. As well as searches on key journals, reference lists, conference proceedings and grey literature were also searched. The search terms were first developed for MEDLINE and then adapted with relevant syntax and subject headings for the other databases. Supplemental digital content 1 shows example of search one and two.

Study selection

Two independent reviewers (SA, EB) assessed studies based on the title and abstract for eligibility. In case of insufficient information, full text articles were retrieved and screened for eligibility. The reviewers discussed findings and reached consensus on eligibility of studies. The percentage agreement between reviewers was estimated using the κ statistic (SPSS for Windows statistical software package IBM SPSS Statistics V.25).

Data extraction

Two reviewers (SA, EB) independently extracted data of eligible studies. Information about study, participants characteristics, outcome measures and measurement properties were extracted. If information was not clear or unavailable in studies, corresponding authors were contacted.

Risk of bias assessment

The risk of bias for each measurement properties was assessed using COSMIN checklist ¹⁴. Adaptions were made for studies of body structure and function e.g. interobserver reliability. This involved removal of inapplicable standards i.e. "was the time interval appropriate?" Each item of measurement property was rated as either 'very good', 'adequate', 'doubtful' or 'inadequate quality' ¹⁴. Subsequently overall methodological quality of measurement property

- was rated based on "the worst score counts principle" 14. Two independent reviewers (SA, EB)
- assessed study quality and inconsistencies were resolved by discussion.

Hypotheses for construct validity and responsiveness

- 75 Hypotheses for evaluating construct validity and responsiveness assessed in included studies,
- were pre-defined ³³ and listed in supplemental digital content 2.

Data analysis and synthesis

- 78 The necessary homogeneity in studies results was insufficient, thus meta-analysis was not
- 79 performed. Results were therefore synthesised and qualitatively summarised ¹⁶. The
- 80 measurement property for each study was rated according to updated criteria for good
- 81 measurement properties as sufficient (+), insufficient (-) or indeterminate (?) ¹⁶. Then,
- 82 evidence was graded using modified Grading of Recommendations Assessment, Development
- and Evaluation (GRADE) approach ¹⁶. Five factors determine quality of evidence: risk of bias,
- inconsistency, indirectness, imprecision and publication bias ³⁴. For evaluating measurement
- properties in systematic reviews of PROM, only four factors were assessed, with fifth factor
- 86 (publication bias) removed ³³.

RESULTS

- 88 The PRISMA flow diagram shows results of both searches, selection process and reasons for
- 89 exclusion (Figure 1).

Search one: inventory of outcome measure

- A list of OM was generated and classified into 28 PROM, 20 PBOM and 10 body structure and
- 92 function OM are listed in supplemental digital content 3. The International Classification of
- 93 Functioning, Disability and Health (ICF) model ²⁵ was used to classify OM into either PBOM
- 94 or body structure and function OM. Agreement between reviewers (SA, EB) for title and

- abstract assessment was excellent (94.0%, Kappa=0.91) and full-text (92.5%, Kappa=0.80).
- The third reviewer (NRH) was consulted twice.

Search two: measurement properties

- There were 16 studies for measurement properties of PROM, 1 study for PBOM and 3 studies
- for body structure and function OM (Table 2). Excellent agreement between reviewers (SA,
- EB) for titles/abstracts (95%, Kappa=0.92) and substantial agreement for full-text articles 13 100
 - (90%, Kappa=0.78) 35. Eleven authors responded from twenty-one who were contacted
- 18 102 clarifying participants age, language of PROM utilized, or for missing data. The third reviewer
 - (NRH) was consulted four times.

Study and outcome measure characteristics

- Detailed information on studies and participant characteristics are shown in Table 2. The OM 26 105
 - included were 9 PROMs (6 disease-specific and 3 generic), 1 PBOM, and 6 body structure and
 - function OM. Detailed description of OMs and their characteristics are shown in Table 3 &
 - Table 4.

Risk of bias

- 39 110 Evaluated measurement properties included, development (n=1), internal consistency (n=3),
 - reliability (n=5), measurement invariance (n=2), measurement error (n=2), hypothesis testing
- for construct validity (n=18), responsiveness (n=2). Results of risk bias assessment are 44 112
 - presented in supplemental digital content 4.

Measurement properties and synthesis of evidence

- Table 5 shows the summary of findings table for results of measurement properties and the
- overall evidence for measurement properties against COSMIN and GRADE approach.

Patient-reported outcome measures

Functional scales of SRS-24 19 displayed sufficient discriminative validity in pre and post-5 6 surgery individuals with AIS ³⁸. While, construct validity of SRS-22 function scale was rated 8 insufficient (Moderate-quality evidence) ^{38,40,41}, and sufficiently responsive ³⁹ (very low-10 122 quality evidence). Measurement invariance of this scale was rated indeterminate since no multiple group factor analysis was performed ⁴³, and the measurement error rated insufficient ⁴². The activity scale of SRS-22r was rated sufficiently reliable as the Interclass Correlation Coefficient (ICC) was 0.76 (0.56–0.80) supported by low-quality evidence. However, internal consistency ¹⁸ was rated indeterminate ³³. The SRS-22r showed insufficient measurement error 20 126 ⁴⁵ (Moderate-quality evidence). A strong correlation between function scale of SRS-22r with mobility scale of Child Health Questionnaire-Child Self-Report Form 87 (CHQ-CF87) (Pearson r=0.73)¹⁸ indicating sufficient convergent validity. Whilst, hypothesis of discriminative validity was not met 44. Thus, evidence for construct validity was downgraded 32 131 for inconsistency. Moreover, the scale was found unresponsiveness to change (low-quality evidence) ⁴⁵. The SRS-30 consists of questions from both SRS-24 and SRS-22. Although no study was identified evaluated its validity or reliability, high-quality evidence indicated that the construct validity of activity scale of SRS-30 was sufficient ⁴⁸. A difference in activity scores (0.50) observed at instrumentations construct before and after surgery 48, whilst measurement invariance was rated indeterminate ⁴⁷. 47 137 Scoliosis Quality of Life Index (SQLI) is a modified version of SRS-22 consisting of physical 52 139 activity domain ³⁶. Very low evidence demonstrated that its content validity is sufficient based on reviewers' ratings only ⁷⁹. The questionnaire was tested for comprehensibility among healthy school children (9.9 years old) only ³⁶. Per COSMIN guidance, those children may not 57 141 consider as representative to population of interest ⁷⁹. The internal consistency of activity scale ⁵⁹ 142

was rated indeterminate, while its reliability was insufficient (ICC =0.46, 0.29 -0.63). The evidence was downgraded due to serious risk of bias and imprecision. Moderate-quality evidence showed that construct validity of this scale was sufficient.

Mobility scale of Patient-Reported Outcomes Measurement Information System (PROMIS) 46 correlated with function scale of SRS-22r (Pearson r=0.65) 46 indicating sufficient construct validity, while functional domains of Paediatrics Outcomes Data Collection Instrument (PODCI) had insufficient construct validity ⁵⁰.

Internal consistency of physical functioning scale of (CHQ-CF87) ¹⁸ rated indeterminate as evidence of sufficient structural validity is not available ³³, while its reliability scale was sufficient (ICC = 0.73, 0.20-0.85) based on low-quality evidence.

The Sport Activity Questionnaire (SAQ) was developed based on a test-retest method, which is considered a reliability study based on COSMIN definitions ¹⁵. A very low-quality evidence showed that reliability of SAQ was sufficient.

In conclusion, according to COSMIN methodology for a PROM to be recommended for use, it should exhibit any level of sufficient content validity and low level of evidence of sufficient internal consistency ³³. None of the identified PROMs in this review met these criteria, thus we are unable to recommend any of these PROMs for use in individuals with AIS. Furthermore, none of these PROM had a high evidence of insufficient measurement properties. Therefore, these PROMs can be used but it require further assessment of the quality of its measurement properties to be recommended for use with individuals with AIS ³³.

Performance-based outcome measure

Timed Up and Go Test (TUG) is the only performance measure identified in this review, its measurement properties tested in AIS. A difference in the time to perform TUG test was found between individuals with AIS having different curve severity ⁵¹, indicating sufficient construct validity ⁵¹.

Body structure and function measures

5

10

52 189

12 173

The Trunk Pelvis Hip Angle (TPHA) test is used to measure mobility of lumbo-pelvic-hip complex⁵⁴. Moderate -quality evidence supported sufficient inter-observer reliability of TPHA $(ICC > 0.942)^{33}$.

Very low evidence showed that criterion validity of Modified Schober Test (MST) 52 rated indeterminate as not all required information reported i.e. amount of correlation with radiographs ⁵². While, its construct validity rated insufficient ⁵².

The construct validity of Fingertip To Floor Test (FTF) and 7th cervical vertebra to Posterior Superior Iliac Spine (C7-PSIS) distance was rated insufficient (moderate-quality evidence) ⁵³. No difference in scores of these tests was found between individuals with mild and severe curves ⁵³. On the other hand, construct validity of Lateral Side Bending (LSB) angle and Axial Rotation was sufficiently different between individuals with severe curves ⁵³.

Interpretability and feasibility

Information about interpretability and feasibility aspects of functional scales included in this review are available in supplemental digital content 5. The majority of these scales had high ceiling effect (20% -44%) and minimal floor effects. An exception to this is physical activity scale of SQLI (minimal ceiling and floor effects) ^{36,37}. The Minimal Clinical Important Difference MCID reported for activity domain for SRS-22 is 0.08 ⁴². While Minimum Detectable Measurement Difference (MDMD) of activity for SRS-22r is 0.24 45. Review studies did not report information about response shift and percentage of missing items. Moreover, limited information found about feasibility aspects. Most of the included PROMs are completed within 2-3 minutes, and it could be concluded that it these PROMs are easy to complete, available in different settings, and available free of charge.

DISCUSSION

52 213

57 215

47 211

This is the first rigorous systematic review identifying OM used to assess physical functioning in individuals with AIS and evaluating their respective measurement properties. Search one enabled generation of a list of OM and search two revealed a few measurement properties studies; comprising nine PROMs, just one PBOM, and six measures of body structure and function. None of the identified PROMs had evidence of sufficient content validity and sufficient internal consistency [34]. Thus, PROMs identified in this review have the potential to be recommended for use but are yet to have the measurement properties investigated. The current evidence showed limited information on the measurement properties of PBOM and body function and structure measure in individuals with AIS.

Patient-reported outcome measure

This review highlights a gap in evidence on content validity of routinely used PROMs that evaluate physical functioning in individuals with AIS. As COSMIN suggested, content validity is the first and most important measurement property to consider when selecting any PROM ⁷⁹. It should be assessed with an interview with both professionals and patients to assess relevance, comprehensiveness, and comprehensibility of items within a PROM ⁷⁹. The identified PROMs lack adequate development process, as many were developed in a population whose mean age was higher than that of individuals with AIS ^{17,19,66,80}. The physical activity scale of SQLI was the only scale where its comprehensibility had been investigated ³⁶, however using healthy children ³⁶ it is not representative of our population of interest ⁷⁹.

The majority of identified measurement properties' studies tested construct validity, which displayed sufficient ratings in most of OMs. Otherwise, internal consistency was undetermined due to lack of evidence of sufficient structural validity. Most of activity scales identified demonstrated high ceiling effects, which affect its ability to assess changes in patient's status

⁵⁹ 216

Performance-based outcome measure

Compared with PROMs just 1 study has investigated measurement properties of a PBOM 51 Where pain 10 and psychological distress 81 may influence the self-reporting of functional ability ¹², it is questionable if PROMs are providing adequate information about actual functional performance of this population. Whilst the TUG test assesses balance, mobility, and walking ability ⁵¹, more evidence-based PBOM are needed to evaluate important and meaningful activities of daily livings for individuals with AIS.

15 223

18 224

Body structure and function measures

23 226

28 228 Radiographs, measured using Cobb angle, are the gold standard measure for evaluating spinal curvature ²⁴. While measurement properties of this measure have been studied before ²⁷, little attention has given to other measures, such as MST and FTF test. These tests are inexpensive, easy, quick measure that does not expose young spines to ionising radiation. When adequate

measurement properties of these OM established, it could serve as a surrogate to radiographs.

This review utilized two-search strategy to enable identification of all types of OM used in

stages. Adherence to the COSMIN methodology as preferred approach for systematic review

of measurement properties is another strength [33]. However, ratings of studies were

determined using lowest score principle, which may underestimate a study's final quality score

[14]. A potential limitation of this review is there are few studies investigating measurement

properties in individuals with AIS, and some that were included where investigating of

Strengths and limitations

AIS. Risk of selection bias was minimized by involving two independent reviewers for all

40 233

45 235

50 237

55 239

60 241

A range of measures are used for physical functioning assessment in individuals with AIS.

CONCLUSION

measurement property was not a primary aim.

The majority of measurement properties studies identified were for PROM with a paucity of

		Adolescent Idiopathic Scoliosis Outcome Measures
1	242	information on PBOM and body structure and function measures. Moreover none of identified
2	243	PROM can be recommended for use in AIS. More measurement properties studies are required
4 5 6	244	to support recommendation of these measures for research and clinical practice.
7 8	245	
9	246	List of Supplemental Digital Contents
11 12	247	Supplemental Digital Content 1. docx
13 14	248	Supplemental Digital Content 2. docx
15 16	249	Supplemental Digital Content 3. docx
17	250	Supplemental Digital Content 4. docx
	251	Supplemental Digital Content 5. docx
20 21		
22 23		
24 25		
25 26		
27		
28 29		
30		
31 32		
33		
34 35		
36		
37 38		
39		
40		
41 42		
43		
44		
45 46		
47		
48		
49		
50 51		
52		
53		
54		

REFERENCES

- 1. Konieczny MR, Senyurt H, Krauspe R. Epidemiology of adolescent idiopathic scoliosis. *J. Child. Orthop.* 2013;7:3-9.
- 2. Weinstein SL, Dolan LA, Cheng JC, et al. Adolescent idiopathic scoliosis. *Lancet* 2008;371:1527-37.
- 3. Hamad A, Ahmed EB, Tsirikos AI. Adolescent idiopathic scoliosis: a comprehensive approach to aetiology, diagnostic assessment and treatment. *J. Orthop. Trauma* 2017;31:343-9.
- 4. Makino T, Kaito T, Kashii M, et al. Low back pain and patient-reported QOL outcomes in patients with adolescent idiopathic scoliosis without corrective surgery. *Springerplus* 2015;4:397.
- 5. Leszczewska J, Czaprowski D, Pawłowska P, et al. Evaluation of the stress level of children with idiopathic scoliosis in relation to the method of treatment and parameters of the deformity. *Sci. World J.* 2012;2012:538409.
- 6. Durmala J, Tomalak W, Kotwicki T. Function of the respiratory system in patients with idiopathic scoliosis: reasons for impairment and methods of evaluation. *Stud Health Technol Inform* 2008;135:237-45.
- 7. Du C, Yu J, Zhang J, et al. Relevant areas of functioning in patients with adolescent idiopathic scoliosis on the International Classification of Functioning, Disability and Health: The patients' perspective. *J. Rehabil. Med.* 2016;48:806-14.
- 8. Dodd S, Clarke M, Becker L, et al. A taxonomy has been developed for outcomes in medical research to help improve knowledge discovery. *J. Clin. Epidemiol.* 2018;96:84-92.

- 9. Tomey KM, Sowers MR. Assessment of physical functioning: a conceptual model encompassing environmental factors and individual compensation strategies. *Phys. Ther.* 2009;89:705-14.
- 10. Bastrom TP, Marks MC, Yaszay B, et al. Prevalence of Postoperative Pain in Adolescent Idiopathic Scoliosis and the Association With Preoperative Pain. *Spine* 2013;38:1848-52.
- 11. LaMontagne LL, Hepworth JT, Cohen F, et al. Adolescent scoliosis: Effects of corrective surgery, cognitive-behavioral interventions, and age on activity outcomes. *Appl. Nurs. Res.* 2004;17:168-77.
- 12. Reiman MP, Manske RC. The assessment of function: How is it measured? A clinical perspective. *J. Man. Manip. Ther.* 2011;19:91-9.
- 13. Bean JF, Olveczky DD, Kiely DK, et al. Performance-based versus patient-reported physical function: what are the underlying predictors? *Phys. Ther.* 2011;91:1804-11.
- 14. Mokkink LB, de Vet HCW, Prinsen CAC, et al. COSMIN Risk of Bias checklist for systematic reviews of Patient-Reported Outcome Measures. *Qual Life Res* 2018;27:1171-9.
- 15. Mokkink LB, Terwee CB, Patrick DL, et al. The COSMIN study reached international consensus on taxonomy, terminology, and definitions of measurement properties for health-related patient-reported outcomes. *J. Clin. Epidemiol.* 2010;63:737-45.
- 16. Prinsen CAC, Mokkink LB, Bouter LM, et al. COSMIN guideline for systematic reviews of patient-reported outcome measures. *Qual. Life Res.* 2018;27:1147-57.
- 17. Asher MA, Min Lai S, Burton DC. Further Development and Validation of the Scoliosis Research Society (SRS) Outcomes Instrument. *Spine* 2000;25:2381-6.
- 18. Glattes RC, Burton DC, Lai SM, et al. The reliability and concurrent validity of the Scoliosis Research Society-22r patient questionnaire compared with the Child Health

- Questionnaire-CF87 patient questionnaire for adolescent spinal deformity. *Spine (Phila Pa 1976)* 2007;32:1778-84.
- 19. Haher TR, Gorup JM, Shin TM, et al. Results of the Scoliosis Research Society instrument for evaluation of surgical outcome in adolescent idiopathic scoliosis. A multicenter study of 244 patients. *Spine (Phila Pa 1976)* 1999;24:1435-40.
- 20. de Kleuver M, Faraj SSA, Holewijn RM, et al. Defining a core outcome set for adolescent and young adult patients with a spinal deformity. *Acta Orthopaedica* 2017;88:612-8.
- 21. Faraj SSA, van Hooff ML, Holewijn RM, et al. Measuring outcomes in adult spinal deformity surgery: a systematic review to identify current strengths, weaknesses and gaps in patient-reported outcome measures. *Eur. Spine J.* 2017;26:2084-93.
- 22. Alamrani S, Rushton A, Gardner A, et al. Outcome measures evaluating physical functioning and their measurement properties in adolescent idiopathic scoliosis: a protocol for a systematic review. *BMJ Open* 2020;10:e034286.
- 23. Moher D, Liberati A, Tetzlaff J, et al. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. *PLoS Med.* 2009;6:e1000097.
- 24. Cobb J. Outline for the study of scoliosis. *Instr Course Lect AAOS* 1948;5:261-75.
- 25. santé Omdl, Organization WH, Staff WHO. *International classification of functioning, disability and health: ICF*ed: World Health Organization, 2001.
- 26. Prowse A, Pope R, Gerdhem P, et al. Reliability and validity of inexpensive and easily administered anthropometric clinical evaluation methods of postural asymmetry measurement in adolescent idiopathic scoliosis: a systematic review. *Eur. Spine J.* 2016;25:450-66.
- 27. Langensiepen S, Semler O, Sobottke R, et al. Measuring procedures to determine the Cobb angle in idiopathic scoliosis: a systematic review. *Eur. Spine J.* 2013;22:2360-71.

- 28. Navarro I, Rosa BND, Candotti CT. Anatomical reference marks, evaluation parameters and reproducibility of surface topography for evaluating the adolescent idiopathic scoliosis: a systematic review with meta-analysis. *Gait Posture* 2019;69:112-20.
- 29. Wade R, Yang H, McKenna C, et al. A systematic review of the clinical effectiveness of EOS 2D/3D X-ray imaging system. *Eur. Spine J.* 2013;22:296-304.
- 30. Fong DY, Lee CF, Cheung KM, et al. A meta-analysis of the clinical effectiveness of school scoliosis screening. *Spine (Phila Pa 1976)* 2010;35:1061-71.
- 31. Wu HD, Liu W, Wong MS. Reliability and validity of lateral curvature assessments using clinical ultrasound for the patients with scoliosis: a systematic review. *Eur. Spine J.* 2020;29:717-25.
- 32. He C, Wong MS. Spinal Flexibility Assessment on the Patients With Adolescent Idiopathic Scoliosis: A Literature Review. *Spine (Phila Pa 1976)* 2018;43:E250-e8.
- 33. Mokkink LB, Prinsen C, Patrick DL, et al. COSMIN methodology for systematic reviews of Patient-Reported outcome measures (PROMs). *User manual* 2018.
- 34. Balshem H, Helfand M, Schünemann HJ, et al. GRADE guidelines: 3. Rating the quality of evidence. *J. Clin. Epidemiol.* 2011;64:401-6.
- 35. Cohen J. A Coefficient of Agreement for Nominal Scales. *Educ. Psychol. Meas.* 1960;20:37-46.
- 36. Feise RJ, Donaldson S, Crowther ER, et al. Construction and validation of the scoliosis quality of life index in adolescent idiopathic scoliosis. *Spine (Phila Pa 1976)* 2005;30:1310-5.
- 37. Parent EC, Hill D, Moreau M, et al. Score distribution of the Scoliosis Quality of Life Index questionnaire in different subgroups of patients with adolescent idiopathic scoliosis. *Spine (Phila Pa 1976)* 2007;32:1767-77.

- 38. Bastrom TP, Bartley C, Marks MC, et al. Postoperative Perfection: Ceiling Effects and Lack of Discrimination With Both SRS-22 and -24 Outcomes Instruments in Patients With Adolescent Idiopathic Scoliosis. *Spine (Phila Pa 1976)* 2015;40:E1323-9.
- 39. Asher M, Min Lai S, Burton D, et al. Scoliosis research society-22 patient questionnaire: responsiveness to change associated with surgical treatment. *Spine (Phila Pa 1976)* 2003;28:70-3.
- 40. Asher M, Min Lai S, Burton D, et al. Discrimination validity of the scoliosis research society-22 patient questionnaire: relationship to idiopathic scoliosis curve pattern and curve size. *Spine (Phila Pa 1976)* 2003;28:74-8.
- 41. Parent EC, Hill D, Mahood J, et al. Discriminative and predictive validity of the scoliosis research society-22 questionnaire in management and curve-severity subgroups of adolescents with idiopathic scoliosis. *Spine (Phila Pa 1976)* 2009;34:2450-7.
- 42. Carreon LY, Sanders JO, Diab M, et al. The minimum clinically important difference in Scoliosis Research Society-22 Appearance, Activity, And Pain domains after surgical correction of adolescent idiopathic scoliosis. *Spine (Phila Pa 1976)* 2010;35:2079-83.
- 43. Verma K, Lonner B, Toombs CS, et al. International utilization of the SRS-22 instrument to assess outcomes in adolescent idiopathic scoliosis: what can we learn from a medical outreach group in Ghana? *J. Pediatr. Orthop.* 2014;34:503-8.
- 44. Berliner JL, Verma K, Lonner BS, et al. Discriminative validity of the Scoliosis Research Society 22 questionnaire among five curve-severity subgroups of adolescents with idiopathic scoliosis. *Spine J* 2013;13:127-33.
- 45. Kelly MP, Lenke LG, Sponseller PD, et al. The minimum detectable measurement difference for the Scoliosis Research Society-22r in adolescent idiopathic scoliosis: a comparison with the minimum clinically important difference. *Spine J* 2019;19:1319-23.

- 46. Fedorak GT, Larkin K, Heflin JA, et al. Pediatric Patient-Reported Outcomes Measurement Information System is Equivalent to Scoliosis Research Society-22 in Assessing Health Status in Adolescent Idiopathic Scoliosis. *Spine* 2019;44:E1206-E10.
- 47. Roberts DW, Savage JW, Schwartz DG, et al. Male-female differences in Scoliosis Research Society-30 scores in adolescent idiopathic scoliosis. *Spine (Phila Pa 1976)* 2011;36:E53-9.
- 48. Lubicky JP, Hanson JE, Riley EH. Instrumentation constructs in pediatric patients undergoing deformity correction correlated with Scoliosis Research Society scores. *Spine* (*Phila Pa 1976*) 2011;36:1692-700.
- 49. Sarwahi V, Wendolowski S, Gecelter R, et al. When Do Patients Return to Physical Activities and Athletics After Scoliosis Surgery?: A Validated Patient Questionnaire Based Study. *Spine (Phila Pa 1976)* 2018;43:167-71.
- 50. Lerman JA, Sullivan E, Haynes RJ. The Pediatric Outcomes Data Collection Instrument (PODCI) and functional assessment in patients with adolescent or juvenile idiopathic scoliosis and congenital scoliosis or kyphosis. *Spine (Phila Pa 1976)* 2002;27:2052-7; discussion 7-8.
- 51. Gao C-C, Chern J-S, Chang C-J, et al. Center of pressure progression patterns during level walking in adolescents with idiopathic scoliosis. *PloS one* 2019;14:e0212161-e.
- 52. Hresko MT, Mesiha M, Richards K, et al. A comparison of methods for measuring spinal motion in female patients with adolescent idiopathic scoliosis. *J. Pediatr. Orthop.* 2006;26:758-63.
- 53. Eyvazov K, Samartzis D, Cheung JPY. The association of lumbar curve magnitude and spinal range of motion in adolescent idiopathic scoliosis: a cross-sectional study. *BMC Musculoskelet. Disord.* 2017;18:51-.

- 54. Stępień A, Guzek K, Pałdyna B, et al. The Trunk-Pelvis-Hip Angle Test is a Reliable Measurement of the Range of the Lower Trunk-Pelvis Rotation in Adolescents. *J Orthop Ther: JORT-1124. DOI* 2018;10:2575-8241.
- 55. Alanay A, Cil A, Berk H, et al. Reliability and validity of adapted Turkish Version of Scoliosis Research Society-22 (SRS-22) questionnaire. *Spine (Phila Pa 1976)* 2005;30:2464-8.
- 56. Monticone M, Carabalona R, Negrini S. Reliability of the Scoliosis Research Society-22 Patient Questionnaire (Italian version) in mild adolescent vertebral deformities. *Eura Medicophys* 2004;40:191-7.
- 57. Bago J, Climent JM, Ey A, et al. The Spanish version of the SRS-22 patient questionnaire for idiopathic scoliosis: transcultural adaptation and reliability analysis. *Spine* (*Phila Pa 1976*) 2004;29:1676-80.
- 58. Hashimoto H, Sase T, Arai Y, et al. Validation of a Japanese version of the Scoliosis Research Society-22 Patient Questionnaire among idiopathic scoliosis patients in Japan. *Spine* (*Phila Pa 1976*) 2007;32:E141-6.
- 59. Cheung KM, Senkoylu A, Alanay A, et al. Reliability and concurrent validity of the adapted Chinese version of Scoliosis Research Society-22 (SRS-22) questionnaire. *Spine* (*Phila Pa 1976*) 2007;32:1141-5.
- 60. Li M, Wang CF, Gu SX, et al. Adapted simplified Chinese (mainland) version of Scoliosis Research Society-22 questionnaire. *Spine (Phila Pa 1976)* 2009;34:1321-4.
- 61. Glowacki M, Misterska E, Laurentowska M, et al. Polish adaptation of scoliosis research society-22 questionnaire. *Spine (Phila Pa 1976)* 2009;34:1060-5.
- 62. Beausejour M, Joncas J, Goulet L, et al. Reliability and validity of adapted French Canadian version of Scoliosis Research Society Outcomes Questionnaire (SRS-22) in Quebec. *Spine (Phila Pa 1976)* 2009;34:623-8.

- 63. Lonjon G, Ilharreborde B, Odent T, et al. Reliability and Validity of the French-Canadian Version of the Scoliosis Research Society 22 Questionnaire in France. *Spine* 2014;39:E26-E34.
- 64. Leelapattana P, Keorochana G, Johnson J, et al. Reliability and validity of an adapted Thai version of the Scoliosis Research Society-22 questionnaire. *J Child Orthop* 2011;5:35-40.
- 65. Adobor RD, Rimeslatten S, Keller A, et al. Repeatability, reliability, and concurrent validity of the scoliosis research society-22 questionnaire and EuroQol in patients with adolescent idiopathic scoliosis. *Spine (Phila Pa 1976)* 2010;35:206-9.
- 66. Asher MA, Lai SM, Glattes RC, et al. Refinement of the SRS-22 Health-Related Quality of Life questionnaire Function domain. *Spine (Phila Pa 1976)* 2006;31:593-7.
- 67. Niemeyer T, Schubert C, Halm HF, et al. Validity and reliability of an adapted german version of scoliosis research society-22 questionnaire. *Spine (Phila Pa 1976)* 2009;34:818-21.
- 68. Antonarakos PD, Katranitsa L, Angelis L, et al. Reliability and validity of the adapted Greek version of scoliosis research society 22 (SRS-22) questionnaire. *Scoliosis* 2009;4:14.
- 69. Schlosser TP, Stadhouder A, Schimmel JJ, et al. Reliability and validity of the adapted Dutch version of the revised Scoliosis Research Society 22-item questionnaire. *Spine J* 2014;14:1663-72.
- 70. Camarini PM, Rosanova GC, Gabriel BS, et al. The Brazilian version of the SRS-22r questionnaire for idiopathic scoliosis. *Braz J Phys Ther* 2013;17:494-505.
- 71. Monticone M, Baiardi P, Calabro D, et al. Development of the Italian version of the revised Scoliosis Research Society-22 Patient Questionnaire, SRS-22r-I: cross-cultural adaptation, factor analysis, reliability, and validity. *Spine (Phila Pa 1976)* 2010;35:E1412-7.

- 72. Sathira-Angkura V, Pithankuakul K, Sakulpipatana S, et al. Validity and reliability of an adapted Thai version of Scoliosis Research Society-22 questionnaire for adolescent idiopathic scoliosis. *Spine (Phila Pa 1976)* 2012;37:783-7.
- 73. Haidar RK, Kassak K, Masrouha K, et al. Reliability and validity of an adapted Arabic version of the Scoliosis Research Society-22r Questionnaire. *Spine (Phila Pa 1976)* 2015;40:E971-7.
- 74. Mousavi SJ, Mobini B, Mehdian H, et al. Reliability and validity of the persian version of the scoliosis research society-22r questionnaire. *Spine (Phila Pa 1976)* 2010;35:784-9.
- 75. Danielsson AJ, Romberg K. Reliability and Validity of the Swedish Version of the Scoliosis Research Society–22 (SRS-22r) Patient Questionnaire for Idiopathic Scoliosis. *Spine* 2013;38:1875-84.
- 76. Scoliosis Research Society web site.
- Available at: http://www.srs.org/professionals/outcomes/srs-30.pdf. SRS-30 Patient Questionnaire.
- 77. Kyrola K, Jarvenpaa S, Ylinen J, et al. Reliability and Validity Study of the Finnish Adaptation of Scoliosis Research Society Questionnaire Version SRS-30. *Spine (Phila Pa 1976)* 2017;42:943-9.
- 78. Carrico G, Meves R, Avanzi O. Cross-cultural adaptation and validity of an adapted Brazilian Portuguese version of Scoliosis Research Society-30 questionnaire. *Spine (Phila Pa 1976)* 2012;37:E60-3.
- 79. Terwee CB, Prinsen CAC, Chiarotto A, et al. COSMIN methodology for evaluating the content validity of patient-reported outcome measures: a Delphi study. *Qual Life Res* 2018;27:1159-70.

- 80. Asher M, Min Lai S, Burton D, et al. The reliability and concurrent validity of the scoliosis research society-22 patient questionnaire for idiopathic scoliosis. *Spine (Phila Pa 1976)* 2003;28:63-9.
- 81. Sanders AE, Andras LM, Iantorno SE, et al. Clinically Significant Psychological and Emotional Distress in 32% of Adolescent Idiopathic Scoliosis Patients. *Spine Deform* 2018;6:435-40.

	Search one	Search two
	(Inventory of outcome measure)	(Measurement properties)
Inclusion criteria	Individuals with AIS (≥10° Cobb angle) ¹ Age 10-18 years old	Individuals with AIS (≥10° Cobb angle) ¹ Age 10-18 years old Mixed cohort studies >50% of participants with AIS
	 Any study design that included assessment of physical functioning for individuals with AIS. No limitations were applied on type of outcome measure, language or location. 	 Measurement properties studies (i.e. content validity, structural validity, construct validity, reliability, and responsiveness) of outcome measure identified in search one.
	Outcome measure defined as following: PROM in form of questionnaires, scales or subscales) designed to evaluate physical functioning in AIS. PBOM, meaning a clinician- observer measure of an "activity" such as the execution of a task or action by an individual ² measured by/or time, or distance. Body structure and function measures defined as "the physiological function of body systems and / or the anatomical parts of body" ^{12,25} .	
Exclusion criteria	Radiographs, laboratory- based measures, anthropometric measures ³⁻⁹ .	 Studies in non-English speaking population Systematic reviews Studies providing normative data Studies providing indirect evidence on measurement properties.

IS indicates Adolescent Idiopathic scoliosis; PROM, Patient Reported Outcome Measure; PBOM, Performance Based Outcome Measure

	Table 2: Studies and participants characteristics								
Patient Re	ported Ou	itcome Mo	easure	T					
Reference	Name of OM	Country	Age (Mean ±SD) Range	Gender (n)	Sample size (n)	Curve type (%), (n)	Curve size Degree ± SD (n)	Type of intervention (n)	Score (mean ± SD)
Feise et al.	SQLI	Canada	14.9 ± 2.4 (10-18)	F(70) M(14)	84	NR	Unbraced 26.1°± 10° Braced 34.3°± 8.7° Postsurgical 31.0°±11.4°	Postsurgical (16) Braced (30) Unbraced (24) Control (14)	81.1± 15.7
Parent et al.	SQLI	Canada	14.7 ±1.9 (8-20)	F(95)	95	Main thoracic (29) Double thoracic (4) Double major (23) Triple major (2) Thoracolumbar/lumbar (20) Thoracolumbar/lumbar, main thoracic (17)	<30° (34) 30°-50° (44) >50° (17)	Surgery	NR
Bastrom et al. 38	SRS- 24,SRS- 22	USA	14.8±2 (10-21)	F(81%)	829	Lenke 1(43%) Lenke 2(20%) Lenke 3 (7%) Lenke 4 (4%) Lenke 5(16%) Lenke 6(10%)	Pre-surgery 55°±13 Post surgery 20°±9	Pre and Post-surgery	Pre-surgery 45°Cobb SRS-22 (4.6±0.5) SRS-24 (4.1±0.5) >80° Cobb SRS-22 (4.2±0.7) SRS-24 (3.8±0.7) Post-surgery <11°Cobb SRS-22(4.6±0.5) SRS-24 (4.17±0.5) >29°Cobb SRS-22 (4.61±0.5) SRS-24 (4.17±0.6)

	1	ı							
Asher et al. Asher et al.	SRS-22 SRS-22	USA	16.4 (10.6 – 47.3)	F(48) M(10)	58 Total (119)	Single (36) Double (19) Triple (3) Thoracic, Thoracolumbar,	63° Largest cobb angle	Surgery Brace,	Function (0 months) 4.1 Function (3 months) 3.3 Function (6 months) 3.9 Function (12 months) 4.2 Function (24 months) 4.3 Control (4.5±0.35) Nonsurgical (4.4±0.36)
			13 (10.7- 15.4) Non-surgical 14 (9.9 -16) Non-surgical untreated 14 (10.8-16) Non-surgical braced 13 (9.9 -15.2) Pre-surgery 14 (10.6-15.8)	(4) Non- surgical F(57) M(11) Non- surgical untreated F(44) M(10) Non- surgical braced F(13) M(1) Pre- surgery F(31) M(1)	Control (19) Nonsurgical (68) Untreated (54) Braced (14) Pre-surgery (32)	Lumbar; double Triple	Non-surgical untreated 27° Braced 31° Pre-surgery 61°	pre-surgery, control	Non-surgical untreated (4.4±0.37) Non-surgical braced (4.5±0.32) Pre-surgery (4.2±0.42)
Parent et al.	SRS-22	Canada	13.5–20 (153) Total (18.6 ± 9.2)	F(153)	153	NR	30° (58) 30°–50°(66) 50° (4)	Observation (107) Brace(32) Pre-surgery (22) Post-surgery (62)	Observation (4.3 ± 0.59) Brace (4.5 ± 0.59) Pre-surgery (4.2 ± 0.58) Post-surgery (4.1 ± 0.60)
Carreon et al. 42	SRS-22	USA	14.3 ± 1.9 (10 –18)	F(735) M(152)	887	NR	53°±18°	Pre & 1 year post-surgery	Pre-surgery 4.15 ± 0.55 Post-surgery 4.23 ± 0.46
Verma et al. 43	SRS-22	USA & Ghana	15.4	F(100) M(60)	160	NR	Ghana 67.2° USA 52°	Pre-surgery	Ghana 3.7 ± 0.8 USA 4.2 ± 0.4
Berliner et al. ⁴⁴	SRS-22r	USA	13.8 (11.0 - 17.2)	F(115) M(40)	155	Non-surgical Thoracic (56.5%) Thoracolumbar (38.7%) Lumbar (4.8%)	Total 43.1° Non-surgical 21.9° Presurgical 57.2°	Non-surgical & pre- surgical	0° -19° (4.5± 0.47) 20° -40° (4.4 ± 0.37) 41° -50° (4.1± 0.69)

(10-22)	F(1,034) 1,281				
	M(247)	Lenke 1(552) Lenke 2 (272) Lenke 3 (93) Lenke 4 (46) Lenke 5 (196) Lenke 6 (120)	NR	1, 2year Post-Surgery	Activity MCID (0.08) MDMD (0.24)
	F(58) Total (70) M(12)	NR	29.8° ± 12.3°	Pre-surgery	SRS-22r (4.5±0.65) CHQ-CF87 (91±15.6)
	F(78.8%) M(21.2%)	Thoracic (67%) Thoracolumbar (21.7%) Lumbar(11.3%)	Thoracic kyphosis °34.1 ±14.9 Lumbar lordosis 54.8°±13.3	Observed, Pre or post- bracing (69.0%) Brace (27.4%) Surgery (3.5%)	PROMIS, Mobility (50.93 ±9.80) SRS-22r, Function (4.5±0.5)
` /	F(83.4%) 744 M(16.5%)	Risser grade M(mean 3.5) F (mean 3.2)	F (53.3°) M (55.9°)	Pre-surgery, 2yr. post- surgery	Pre-surgery F (4.2) M (4.2) Post-surgery F (4.3) M (4.4)
USA 15.6 ±1.7	F(75%) 356	NR	NR	Pre-surgery, 2yr. post- surgery	Pre-surgery (4.18 ± 0.55) Post-surgery (4.34 ± 0.51)
USA 15 (13 –17)	F(71) 95 M(24)	NR	Pre-surgery 51.08° Post-surgery 15.98°	NR	NR
America 15.2 (11.7– 18.8) Patient 15.3 (11.7– 20.9)	Parent F(88) M(9) Patient F(86) M(9)	Thoracic (17) Thoracolumbar (6) Lumbar(7) Double curve (17)	10-29° (n=23) 30-49° (n=20) >50° (n=4)	1 year Post surgery	Upper extremity (96.8± 9.9) Transfer (97.6± 4.7) Sport & Physical Function (85.5±17.5) Global function (89.4±9.8)
Measure					
USA Mild AIS 14.9 ± 1.7 Moderate AIS 16.4 ± 3.3 Severe AIS 15.3 ± 3.1	NR AIS (30) Control (30)	Right-sided Thoracolumbar	Mild AIS 19.9°±4.3 Moderate AIS 31.8°±4.2 Severe AIS 53.4°±16.1	Pre-treatment	TUG (Seconds) Mild (6.8±1.5) Moderate (6.9±0.9) Severe (6.5±0.8) Healthy control (6.0±0.6)
	Moderate AIS 16.4±3.3 Severe AIS	Moderate AIS 16.4±3.3 Severe AIS	Moderate AIS 16.4±3.3 Severe AIS	Moderate AIS 16.4±3.3 Severe AIS Severe AIS	Moderate AIS 16.4±3.3 Severe AIS Severe AIS 53.4°±16.1

Hresko et al. 52	MST	USA	14.2 ± 1.9 (11.3-18.6)	F(37)	37	Thoracic Lumbar	Thoracic 40°±20° Lumbar 31°±12°	Pre-treatment	5.7 ± 2.2 cm
Eyvazov et al. 53	MST FTF test, Axial rotation, LSB, ΔC7-PSIS	China	15.7 ± 4.1	M(12) F(46)	58	Lenke 5 (Thoracolumbar/ lumbar)	Group A 25° ± 7.1° Group B 49.8° ± 13.6° Tot 34°± 9.2°	Pre-treatment	Modified Schober's (cm) Group A: (20.6 ± 1.4) Group B: (20.3 ± 1.2) FTF test (cm) Group A: (10.1 ± 11.2) Group B: (11 ± 10.3) Δ C7-PSIS $(27.6 \pm 1.8\%)$ LSB (degrees) Group A: (66.6 ± 13.4) Group B: (57.8 ± 14.3) Axial rotation (degrees) Group A: (90.1 ± 21.9) Group B: (5.9 ± 19.6)
Stepien et al. ⁵⁴	TPHA test	Poland	AIS (12.7 ± 2.6) Control (11.8 ±2.5)	F(98)	Control (49) AIS (49)	Risser sign Grade 0 (14) Grade 1 (11) Grade 2 (6) Grade 3 (3) Grade 4 (9) Grade 5 (6)	Thoracic 27.7° ±13.4° Lumbar 25.8°±10.5°	Physiotherapy	AIS Left TPHA -10.93°±4.64° Right TPHA -2.37°± 8.30° Control Left TPHA -11°± 3.30° Right TPHA -8.64°±4.70°

AIS indicates Adolescent Idiopathic scoliosis; CHQ-CF87, Child Health Questionnaire- Child Self-Report Form 87; C7-PSIS, Cervical 7 to Posterior Superior Iliac Spine; F, Female; FTF, Fingertip To Floor Test; LSB, Lateral Side Bending; M, Male; MCID, Minimal Clinically Important Difference; MDMD, Minimal Difference; MST, Modfied Schober Test; NR, Not Reported; OM, Outcome Measure; PODCI, Paediatrics Outcomes Data Collection Instrument; PROMIS, Patient-Reported Outcomes Measurement Information System; SAQ, Sport Activity Questionnaire; SD, Standard Deviation; SRS, Scoliosis Research Society; SRS-22r, Scoliosis Research Society-22Revised; SQLI, Scoliosis Quality of Life Index; TPHA, Trunk Pelvis Hip Angle test; TUG, Timed Up and Go Test; USA, United State of America.

Table 3:	Table 3: Patient-reported outcome measures characteristics								
PROMS	Country	Sub-scale items(n)	Target population	Mode of administration	Recall period	Response options	Scoring system	Available translations	
SRS-24 19	USA	General Function (3) Function after surgery (2) Function-activity (3)	AIS	Self- administrated	Now, post- surgery	5 response options	1-5	-	
SRS-22 ³⁹	USA	Function/Activity (5)	AIS	Self- administrated	Now, post- surgery	5 response options	1-5	Turkish ⁵⁵ , Italian ⁵⁶ , Spanish ⁵⁷ , Japanese ⁵⁸ , Traditional Chinese ⁵⁹ , Simplified Chinese ⁶⁰ , Polish ⁶¹ , French ^{62,63} , Thai ⁶⁴ , Norwegian ⁶⁵	
SRS-22r 66	USA	Function/Activity (5)	AIS	Self- administrated	Now, post- surgery	5 response options	1-5	German ⁶⁷ , Greek ⁶⁸ , Dutch ⁶⁹ , Chinese ⁵⁹ , Brazilian ⁷⁰ , Italian ⁷¹ , Thai ⁷² , Arabic ⁷³ , Persian ⁷⁴ , Swedish ⁷⁵	
SRS-30 ⁷⁶	USA	Function/Activity (5) post-surgery questions (2)	AIS	Self- administrated	Now, post- surgery	Function/Activity (5 response options) Post-surgery (3 response options)	Function (1-5) post-surgery (1-3)	Finnish ⁷⁷ Brazilian ⁷⁸	
CHQ- CF87 ¹⁸	USA	Physical Functioning (9)	Generic	Self- administrated	NR	4, 5, 6 Response options	0-100	-	
SQLI 36	Canada	Physical activity (5)	AIS	Self- administrated	Four weeks	5 Response options	0-4	-	
SAQ ⁴⁹	USA	Total (24) School, Gym, Carry backpack, Bend over, Running	AIS	Self- administrated	Post- surgery	NR	NR	-	
PROMIS	USA	Mobility	Generic	Self- administrated	7-day	5 response options	Mean T-score 50, SD 10	-	
PODCI 50	North America	Upper Extremity Functioning, Transfers& basic Mobility Sport & Physical Function Global function	Generic Paediatric orthopaedic conditions	Self-administrated Parent-report Adolescents report	NR	3-6	0-100	-	

AIS indicates Adolescent Idiopathic scoliosis; CHQ-CF87, Child Health Questionnaire: SD, Standard Deviation; SRS, Scoliosis Research Society: SRS-22r, Scoliosis Research Society: 22revised; SQLI, Scoliosis Quality of Life Index; USA, United State of America.

Outcome measure (Reference)	Activity	Required Equipment	Number of trials	Parameter measured
TUG 51	Stand from chair, walk 3m, return, sit down	Chair, stopwatch, walking space	3 trials	Average of time in seconds
MS Test ^{52,53}	Marks on PSIS, keep knees straight, bend forward and touch the floor	Tape measure	2-3 trials	Average of distance is cm
FTF test ⁵³	Stood upright, bend forward and touch the floor	Tape measure	2 trials	Average of distance is cm
C7-PSIS distance ⁵³	Stand upright, maximally flex and extend neck, distance measured between C7 spinous process and PSIS	Tape measure	2 trials	Average of distance in cm
LSB angles ⁵³	In upright posture, knees straight, bend to the side without rotation	Goniometer	2 trials	Average angle in degrees between line joining PSIS and C7
Axial rotation 53	Seated position, locked both arms in front of body with fixed pelvic, shoulder rotation controlled by a goniometer holder device	Goniometer	2 trials on left and right side	Average angle in degrees
TPHA ⁵⁴	Supine, flex & pull lower limbs, then move limbs to the left or right side	Plurimeter	Three times on each side of body	Average of angle in degrees

C7-PSIS indicates Cervical 7 to Posterior Superior Iliac Spine; FTF, Fingertip To Floor Test; LSB, Lateral Side Bending; MST, Modfied Schober Test; PBOM, Performance Based Outcome Measure; TPHA, Trunk Pelvis Hip Angle test; TUG, Timed Up and Go test.

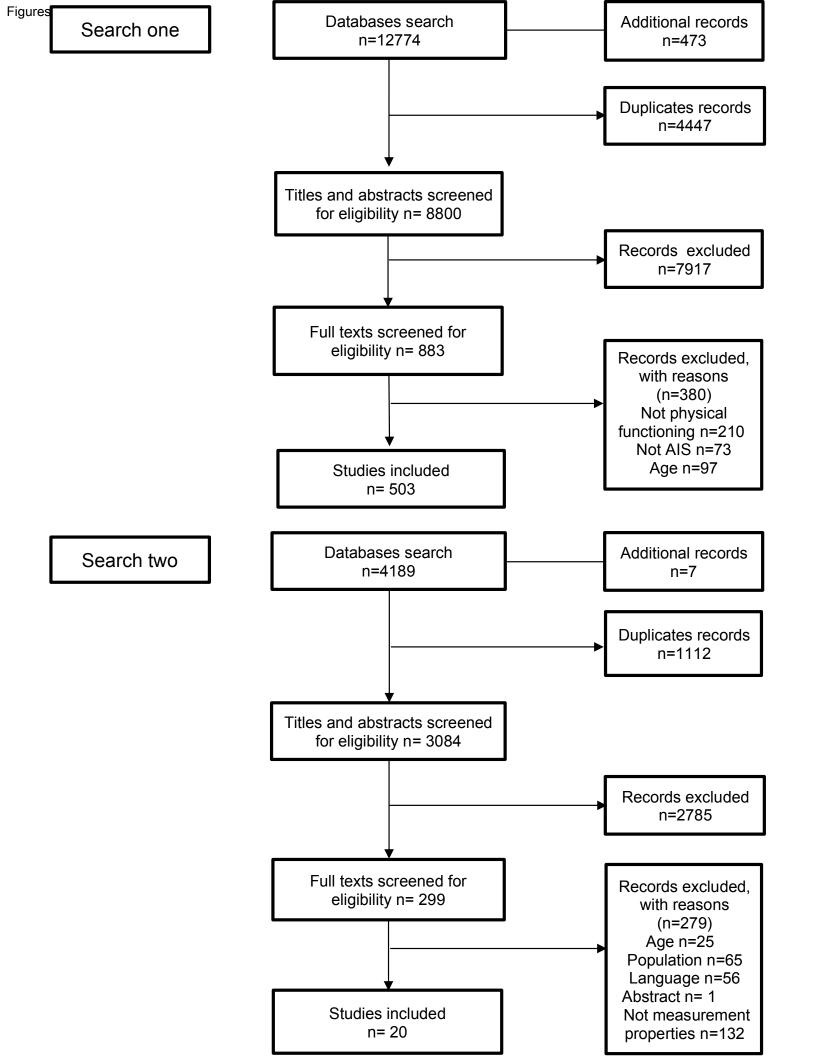
able 5: Summary of finding	ngs table for the measurement properties	s of outcome measure		
Measurement property	Outcome measure (Subscale)	Summary result	Overall rating	Quality of evidence
	SRS-22r (Activity)	∝ = 0.82	?	Moderate (Imprecision)
Internal consistency	SQLI (Physical activity)	∝ = 0.82 (0.76–0.88)	?	Moderate (Imprecision)
	CHQ-CF87 (Physical function)	∝ = 0.89	?	Moderate (Imprecision)
	SRS-22r (Activity)	ICC=0.76 (0.56- 0.80)	+	low (One study adequate quality, Imprecision)
	SQLI (Physical activity)	ICC= 0.46 (0.29 -0.63)	-	Low (One study adequate quality, Imprecision)
Reliability	CHQ-CF87 (Physical function)	ICC=0.73 (0.20- 0.85)	+	Low (One study adequate quality, Imprecision)
	SAQ	Kappa k ≥ 0.70	+	Very low (One study of doubtful quality)
	TPHA Test	ICC= 0.85 (0.95-0.98)	+	Moderate (Imprecision)
Cross- cultural validity\	SRS-22 (Activity)	No multiple group factor analysis performed	?	Very low (one study inadequate quality, Imprecision
measurement invariance	SRS-30 (Function/Activity)	No multiple group factor analysis performed	?	Moderate (one study adequate quality)
Measurement error	SRS-22 (Activity)	SDC (0.24) >MIC(0.08)	-	Moderate (one study of adequate quality)
	SRS-22r	SDC (0.41)> MIC(.08)	-	Moderate (one study of adequate quality)

	SRS-24 (Function)	2 hypotheses confirmed	+	High (One study very good quality
	SRS-22 (Activity)	2 out of 9 hypotheses confirmed	-	Moderate (Inconsistency)
	SRS-22r (Function)	4 hypotheses confirmed	+	Moderate (Inconsistency)
	SRS-30 (Function)	1 hypothesis confirmed	+	High (One study very good quality)
Construct validity	SQLI (Physical activity)	2 hypotheses confirmed	+	Moderate (Imprecision)
Construct valuaty	PODCI (functional scales)	2 hypotheses out of 5 confirmed	-	Moderate (One study adequate quality)
	PROMIS (Mobility)	1 hypothesis confirmed +		Moderate (One study adequate quality)
	TUG test	2 hypotheses out of 3 confirmed	+	Moderate (One study adequate quality)
	MST, FTF Test, C7-PSIS	3 hypotheses not confirmed	_	Moderate (Imprecision)
	LSB ,Axial rotation	2 hypotheses confirmed	+	Moderate (Imprecision)
Criterion validity	MST	Not all information for '+' reported	?	Moderate
Pagnongiyanaga	SRS-22 (Activity)	4 hypotheses confirmed	+	Very low (One study of doubtful quality, Imprecision)
Responsiveness	SRS-22r (Function)	1 hypothesis not confirmed	_	Low (One study doubtful quality)
	·	•	•	•

CHQ-CF87 indicates Child Health Questionnaire- Child Self-Report Form 87; C7-PSIS, Cervical 7 to Posterior Superior Iliac Spine; FTF, Fingertip To Floor Test; ICC, Interclass Correlation Coefficient; LSB, Lateral Side Bending; MIC, Minimal Important Change; MST, Modfied Schober Test, PODCI: Paeditatrics Outcomes Data Collection Instrument,
PROMIS: Patient-Reported Outcomes Measurement Information System, SAQ: Sport Activity Questionnaire, SDC:Small Detectable Change, SRS: Scoliosis Research Society, SRS-22r: Scoliosis Research Society-22Revised, SQLI: Scoliosis Quality of Life Index, TPHA: Trunk Pelvis Hip Angle test, TUG: Timed Up and Go, CCronbach alpha, + =
Sufficient,? = Indeterminate, - = Insufficient.

Figures legends

Figure 1: PRISMA flow diagram of both searches and selection process



Supplemental Data File 1 (doc.)

Click here to access/download

Supplemental Data File (doc., pdf., xls., etc.)

S1.docx

Supplemental Data File 2 (doc.)

Click here to access/download

Supplemental Data File (doc., pdf., xls., etc.)

S 2.docx

Supplemental Data File 3 (doc.)

Click here to access/download

Supplemental Data File (doc., pdf., xls., etc.)

\$3.docx\$

Supplemental Data File 4 (doc.)

Click here to access/download

Supplemental Data File (doc., pdf., xls., etc.)

S 4.docx

Supplemental Data File 5 (doc.)

Click here to access/download

Supplemental Data File (doc., pdf., xls., etc.)

S 5.docx

Physical functioning in adolescents with idiopathic scoliosis: A systematic review of outcome measures and their measurement properties.

- Two searched-strategy performed on all types of outcome measure used in physical functioning assessment for Adolescent with Idiopathic Scoliosis (AIS).
- Most of studies of measurement properties were evaluating Patient Reported Outcome Measure (PROM) with paucity of information on Performance-Based Outcome Measure (PBOM), and body structure and function outcome measures.
- Based on COSMIN methodology, none of measure identified in this review can be recommended for use in individuals with AIS.