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OMERACT Core outcome measurement set for shared decision making in rheumatic and musculoskeletal conditions: a scoping review to identify candidate instruments

Florian Naye ^a, Karine Toupin-April ^{b,c,d,e}, Maarten de Wit ^f, Annie LeBlanc ^{g,h}, Olivia Dubois ^a, Annelies Boonen ¹, Jennifer L. Barton ^j, Liana Fraenkel ^k, Linda C. Li ^l, Dawn Stacey ^{m,n}, Lyn March ^{o,p}, Claire E.H. Barber ^q, Glen Stewart Hazlewood ^r, Francis Guillemin ^s, Susan J. Bartlett ^{t,u,v}, Dorthe B. Berthelsen ^w, Kate Mather ^x, Laurent Arnaud ^y, Akpabio Akpabio ^z, Adewale Adebajo ^{aa}, Grayson Schultz ^{ab}, Victor S. Sloan ^{ac,ad}, Tiffany K. Gill ^{ae}, Saurab Sharma ^{af,ag}, Marieke Scholte-Voshaar ^{ah,ai}, Francesco Caso ^{aj}, Elena Nikiphorou ^{ak,al}, Samah Ismail Nasef ^{am}, Willemina Campbell ^{an}, Alexa Meara ^{ao}, Robin Christensen ^{ap}, Maria E. Suarez-Almazor ^{aq}, Janet Elizabeth Jull ^{ar}, Rieke Alten ^{as}, Esi M. Morgan ^{at}, Yasser El-Miedany ^{au}, Jasvinder A. Singh ^{av}, Jennifer Burt ^{aw}, Arundathi Jayatilleke ^{ax}, Ihsane Hmamouchi ^{ay}, Francisco J. Blanco ^{az}, Anthony P. Fernandez ^{ba}, Sarah Mackie ^{bb}, Allyson Jones ^{bc}, Vibeke Strand ^{bd}, Sara Monti ^{be}, Simon R. Stones ^{bf}, Rebecca R. Lee ^{bg,bh}, Sabrina Mai Nielsen ^{bi}, Vicki Evans ^{bj}, Hemalatha Srinivasalu ^{bk,bl}, Thomas Gérard ^a, Juliette LeBlanc Demers ^g, Roxanne Bouchard ^g, Théo Stefan ^g, Michèle Dugas ^g, Frédéric Bergeron ^{bm}, Dorcas Beaton ^{bn}, Lara J. Maxwell ^{bo}, Peter Tugwell ^{bp}, Simon Décary ^{a,*}

- ^a Faculty of Medicine and Health Sciences, School of Rehabilitation, Research Centre of the CHUS, CIUSSS de l'Estrie-CHUS, Université de Sherbrooke, 3001, 12e Avenue Nord. Sherbrooke. Ouebec J1H 5N4. Canada
- ^b School of Rehabilitation Sciences, Faculty of Health Sciences, University of Ottawa, Ottawa, Canada
- ^c Department of Pediatrics, Faculty of Medicine, University of Ottawa, Ottawa, Canada
- ^d Children's Hospital of Eastern Ontario Research Institute, Ottawa, Canada
- ^e Institut du savoir Montfort, Ottawa, Canada
- f Patient Research Partner, Amsterdam, The Netherlands
- g Department of Family Medicine and Emergency Medicine, Université Laval, Quebec City, Canada
- h VITAM Centre de recherche en santé durable, Quebec City, Canada
- i Department of Internal Medicine, Division of Rheumatology, Maastricht University Medical Center and Caphri Research Institute, Maastricht University, Maastricht, The Netherlands
- ^j VA Portland Health Care System, Oregon Health & Science University, Portland, USA
- k Department of Internal Medicine, Yale University, New Haven, USA
- ¹ Department of Physical Therapy, Arthritis Research Canada, University of British Columbia, Vancouver, Canada
- ^m School of Nursing, University of Ottawa, Ottawa, Canada
- ⁿ The Ottawa Hospital Research Institute, Ottawa, Canada
- O Department of Medicine, The University of Sydney, Sydney, Australia
- P Institute of Bone and Joint Research, Department of Rheumatology, Royal North Shore Hospital, Sydney, Australia
- q Department of Medicine, Department of Community Health Sciences, Cumming School of Medicine, University of Calgary, Calgary, Canada
- ^r Department of Medicine, University of Calgary, Calgary, Canada
- s INSPIIRE, Université de Lorraine, Inserm, Nancy, France
- t Divisions of Clinical Epidemiology, Rheumatology and Respiratory Epidemiology and Clinical Trials Unit, McGill University, Canada
- ^u Research Institute McGill University Health Centre, Canada
- ^v Johns Hopkins Medicine Division of Rheumatology, Montreal, Canada
- ** Section for Biostatistics and Evidence-Based Research, The Parker Institute, Bispebjerg and Frederiksberg Hospital, Copenhagen & Research Unit of Rheumatology, Department of Clinical Research, Odense & Department of Rehabilitation, Municipality of Guldborgsund, Odense University Hospital, University of Southern Denmark, Nykoebing, Denmark

https://doi.org/10.1016/j.semarthrit.2023.152344

^{*} Corresponding author.

E-mail address: Simon.decary@usherbrooke.ca (S. Décary).

- ^x Patient Research Partner, Toronto, Canada
- y Department of Rheumatology, CRMR RESO, University Hospitals of Strasbourg, France
- ² Royal National Hospital for Rheumatic Diseases, Bath, UK
- ^{aa} Faculty of Medicine, Dentistry and Health, University of Sheffield, UK
- ^{ab} Patient Research Partner, Ohio, USA
- ac Sheng Consulting LLC, Flemington, NJ, USA
- ad The Peace Corps, Washington, DC, USA
- ^{ae} Faculty of Health and Medical Sciences, Adelaide Medical School, The University of Adelaide, Australia
- ^{af} School of Health Sciences, Faculty of Medicine and Health, University of New South Wales, Sydney, Australia
- ^{ag} Centre for Pain IMPACT, Neuroscience Research Australia, Sydney, Australia
- ah Patient Research Partner, Department of Pharmacy and Department of Research & Innovation, Sint Maartenskliniek, Nijmegen, The Netherlands
- ai Department of Pharmacy, Radboud university medical center, Nijmegen
- ^{aj} Department of Clinical Medicine and Surgery, University of Naples Federico II, Italy
- ak Centre for Rheumatic Diseases, King's College Hospital, School of Immunology and Microbial Sciences, King's College London, UK
- al Rheumatology Department, King's College Hospital, London, UK
- am Department of Rheumatology and Rehabilitation, Faculty of Medicine, Suez Canal University, Ismailia, Egypt
- ^{an} Patient research partner, Toronto Western Hospital, University Health Network, Canada
- ^{ao} Division of Rheumatology, The Ohio State University, Columbus, USA
- ^{ap} Musculoskeletal Statistics Unit, The Parker Institute, Bispebjerg and Frederiksberg Hospital, Copenhagen, & Department of Rheumatology, Odense University Hospital, Denmark
- aq Department of General Internal Medicine, Section of Rheumatology and Clinical Immunology, University of Texas MD Anderson Cancer Center, Houston, USA
- ar School of Rehabilitation Therapy, Queen's University, Kingston, Canada
- as Department of Internal Medicine II, Rheumatology Research Center, Rheumatology, Clinical Immunology, Osteology, Physical Therapy and Sports Medicine, Schlosspark-Klinik, Charité, University Medicine Berlin, Berlin, Germany
- at Department of Pediatrics, University of Washington, Division of Rheumatology, Seattle Children's Hospital, Seattle, Washington, USA
- ^{au} Canterbury Christ Church University, King's College London, UK
- av University of Alabama at Birmingham, Birmingham, USA
- ^{aw} Newfoundland and Labrador Health Services, St. Clare's Mercy Hospital, St John's, Newfoundland and Labrador, Canada
- ^{ax} Lewis Katz School of Medicine, Temple University, Philadelphia, USA
- ^{ay} Health Sciences Research Centre (CReSS), Faculty of Medicine, International University of Rabat (UIR), Rabat, Morocco
- ^{az} Departamento de Fisioterapia, Medicina y Ciencias Médicas, Universidad de A Coruña, A Coruña, Spain
- ba Departments of Dermatology and Pathology, Cleveland Clinic, Cleveland, Ohio, USA
- bb Leeds Institute of Rheumatic and Musculoskeletal Medicine, Chapel Allerton Hospital, University of Leeds, Leeds Teaching Hospitals NHS Trust, Leeds, UK
- ^{bc} Department of Physical Therapy, Faculty of Rehabilitation Medicine, University of Alberta, Edmonton, Canada
- bd Division of Immunology/Rheumatology, Stanford University, Stanford, California, USA
- be Department of Rheumatology, Policlinico S. Matteo, IRCCS Fondazione, University of Pavia, Pavia, Italy
- ^{bf} Patient research partner, Envision Pharma Group, Wilmslow, UK
- bg Centre for Epidemiology Versus Arthritis, Centre for Musculoskeletal Research, Division of Musculoskeletal and Dermatological Sciences, Faculty of Biology, Medicine and Health, Manchester Academic Health Science Centre, University of Manchester, UK
- bh National Institute for Health Research Biomedical Research Centre, Manchester University Hospital NHS Trust, Manchester, UK
- bi Musculoskeletal Statistics Unit, The Parker Institute, Department of Rheumatology, Odense University Hospital, and University of Southern Denmark, Copenhagen, Denmark, Copenhagen, Denmark
- bj Patient Research Partner and Discipline of Optometry, Faculty of Health, University of Canberra, Canberra, Australia
- bk Pediatric Rheumatology, Children's National Hospital, Washington DC, USA
- bl GW School of Medicine, Washington DC, USA
- bm Université Laval, Quebec City, Canada
- bn Institute for Work & Health, Toronto, Canada
- bo Centre for Practice Changing Research, Ottawa Hospital Research Institute and Faculty of Medicine, University of Ottawa, Ottawa, Canada
- bp Division of Rheumatology, Department of Medicine, and School of Epidemiology and Public Health, Faculty of Medicine, University of Ottawa; Clinical Epidemiology Program. Ottawa Hospital Research Institute. Ottawa. Canada

ARTICLE INFO

Keywords: Shared decision making Rheumatic diseases Musculoskeletal diseases Measurement instruments Core outcome measurement set

ABSTRACT

Objectives: Shared decision making (SDM) is a central tenet in rheumatic and musculoskeletal care. The lack of standardization regarding SDM instruments and outcomes in clinical trials threatens the comparative effectiveness of interventions. The Outcome Measures in Rheumatology (OMERACT) SDM Working Group is developing a Core Outcome Set for trials of SDM interventions in rheumatology and musculoskeletal health. The working group reached consensus on a Core Outcome Domain Set in 2020. The next step is to develop a Core Outcome Measurement Set through the OMERACT Filter 2.2.

Methods: We conducted a scoping review (PRISMA-ScR) to identify candidate instruments for the OMERACT Filter 2.2 We systematically reviewed five databases (Ovid MEDLINE®, Embase, Cochrane Library, CINAHL and Web of Science). An information specialist designed search strategies to identify all measurement instruments used in SDM studies in adults or children living with rheumatic or musculoskeletal diseases or their important others. Paired reviewers independently screened titles, abstracts, and full text articles. We extracted characteristics of all candidate instruments (e.g., measured construct, measurement properties). We classified candidate instruments and summarized evidence gaps with an adapted version of the Summary of Measurement Properties (SOMP) table.

Results: We found 14,464 citations, read 239 full text articles, and included 99 eligible studies. We identified 220 potential candidate instruments. The five most used measurement instruments were the Decisional Conflict Scale (traditional and low literacy versions) (n=38), the Hip/Knee-Decision Quality Instrument (n=20), the Decision Regret Scale (n=9), the Preparation for Decision Making Scale (n=8), and the CollaboRATE (n=8). Only 44 candidate instruments (20%) had any measurement properties reported by the included studies. Of these instruments, only 57% matched with at least one of the 7-criteria adapted SOMP table.

Conclusion: We identified 220 candidate instruments used in the SDM literature amongst people with rheumatic and musculoskeletal diseases. Our classification of instruments showed evidence gaps and inconsistent reporting

of measurement properties. The next steps for the OMERACT SDM Working Group are to match candidate instruments with Core Domains, assess feasibility and review validation studies of measurement instruments in rheumatic diseases or other conditions. Development and validation of new instruments may be required for some Core Domains.

Abbreviations

CROM clinician reported outcome measure
DOI decision Quality Instruments

FAPI Fragebogen zur arzt-patienten-interaktion
MASRI medication adherence self-report inventory

N/A not applicable NR not reported OA osteoarthritis

OMERACT outcome measures in rheumatology

Option scale Observing Patient Involvement in Decision Making

instrument

PCC population, concept and context

PRESS peer review of electronic search strategies

PRISMA preferred reporting items for systematic reviews and meta-

analysis

PRISMA-ScR preferred reporting items for systematic reviews and

meta-analysis extension for scoping reviews

PROM patient reported outcome measure RMDs rheumatic and musculoskeletal diseases

SD standard deviation SDM share decision making

SOMP summary of measurement properties

Introduction

Shared decision making (SDM) is a process by which clinicians collaborate with patients to provide high-quality care based on best evidence and the patient's needs, values, and preferences [1,2]. Two Cochrane systematic reviews of SDM interventions reported inconsistency in the impact of SDM on decision making outcomes across trials, as well as heterogeneity of measurement instruments used to assess specific outcomes [3–5]. This lack of standardization is a significant threat to comparative effectiveness research and could adversely affect the conclusions of systematic reviews [6]. To address these inconsistencies, the Outcome Measures in Rheumatology (OMERACT) SDM Working Group (WG) is developing a Core Outcome Set for trials of SDM interventions in rheumatic and musculoskeletal diseases (RMDs) (https://omeract.org/working-groups/sdm/).

OMERACT is an independent international initiative of researchers, clinicians and patients that is at the forefront of Core Outcome Set development [7]. A Core Outcome Set is defined as an agreed minimal standardized set of outcomes measures, which should be used and reported as a minimum in all clinical trials on a specific area [8,9]. OMERACT uses a rigorous stepwise approach to developing Core Outcome Sets including 1) determining a Core Domains Set that should be measured in all randomized controlled trials and longitudinal observational studies (i.e, what to measure in terms of outcomes, also called domains) [10] and 2) determining the Core Outcome Measurement Set (i.e., how to measure the domains) [11,12].

The OMERACT SDM WG is classified as a 'bolt-on' group. 'Bolt-on groups' describe the additional domains and instruments that are part of a specific intervention, and which are measured in addition to disease-specific core outcome sets. In a clinical trial of SDM interventions, the trial must measure both the core outcome set specific to the concept of SDM and include the disease-specific core outcome set of the clinical trial's study population. By doing so, we ensure that we measure both intervention-specific and disease-specific outcomes.

The OMERACT SDM WG conducted literature reviews, surveys,

interviews, and consensus meetings to develop the Core Domains Set for SDM [13–15]. In 2020, the OMERACT SDM Working Group reached consensus on the Core Domains Set to use in rheumatology and musculoskeletal trials of SDM interventions through virtual consensus meetings with 149 patients, caregivers, clinicians, and researchers [16]. The definitions of the domains include 1) knowledge of options including their potential benefits and harms, 2) chosen option aligned with each study participant's values, 3) certainty in the chosen option, 4) satisfaction with the decision making process, 5) adherence to the chosen option, and 6) potential negative consequences of the SDM intervention [16]. The next step in the OMERACT process is to identify candidate instruments to assess the Core Domains and then, to determine the Core Outcome Measurement Set using the OMERACT Filter 2.2¹². To our best knowledge, no study to date has identified candidate instruments to assess one or more of the six Core Domains for SDM in RMDs.

Objectives

The primary objective of this scoping review was to identify all available measurement instruments reported in the SDM literature for people with RMDs. A secondary objective was to explore the content of the included studies to identify evidence gaps for the future application of the OMERACT Filter 2.2 (i.e., measurement properties, domain match, and feasibility).

Methods

Research questions and OMERACT Filter 2.2 framework

This scoping review addresses the following research question: What are candidate instruments for SDM outcomes amongst people with RMDs? We define "measurement instrument" as a tool that is used to measure a quality or quantity of a variable. *Boers* et al. define this as "tool may be a single question, a questionnaire, a score obtained through physical examination, a laboratory measurement, a score obtained through observation of an image, and so on" [17]. Data from this scoping review informs the identification of candidate measurement instruments for each of the Core Domains. These candidate instruments will then follow the methodological framework of the OMERACT Filter 2.2.

The OMERACT Filter 2.2¹¹ is an instrument selection algorithm to determine a Core Outcome Measurement Set. This filter relies on three pillars of evidence to ensure that a measurement instrument can be included in a Core Outcome Measurement Set¹²: 1) Truth (i.e., domain matching and construct validity), 2) Discrimination (i.e., test-retest reliability, longitudinal construct validity, clinical trial dissemination, thresholds of meaning), and 3) Feasibility. A first working group consensus is based on domain match and feasibility to determine the best candidate measurement instruments [11]. A second working group consensus is organized after a critical appraisal of the measurement properties of these best candidate instruments to select those to integrate in the Core Outcome Measurement Set [11,12]. This scoping review is a preliminary phase to identify evidence gaps prior to answer the three Pillars of the Filter 2.2.

Study design

We conducted a scoping review based on the current update of the

Joanna Briggs Institute guidance [18]. We reported our findings according to the extension for scoping review of the Preferred Reporting Items for Systematic Reviews and Meta-Analysis (PRISMA-ScR). We registered the protocol in Open Science Framework (doi.org/10.1760 5/OSF.IO/4T26R).

Selection and eligibility criteria

Table 1 presents the study eligibility criteria according to the Population, Concept and Context (PCC) framework [18].

Literature search strategy

An information specialist (FB) in consultation with the review team, developed and tested the search strategies using an iterative process. The MEDLINE strategy was peer-reviewed by another senior information specialist using the Peer Review of Electronic Search Strategies (PRESS) checklist [19]. Strategies utilized controlled vocabulary and key words to operationalize the population and concept of our PCC framework (Table 3). We adjusted vocabulary and syntax across databases. There were no date restrictions. Specific details regarding the search strategies for each database are in Supplemental material.

We undertook the systematic search using five databases: MED-LINE® via Ovid, Embase via Ovid, Cochrane Library databases via Ovid, CINAHL via EBSCO and Web of Science. All searches were performed on March 3rd, 2023. We conducted a structured handsearching of primary studies with backward (i.e., inspecting the references that are cited in the included study [20]) and forward (i.e., using a citation index to identify studies that cite the included study [20]) citations of all the included studies on Web of Science [20]. We used literature reviews (e. g., Cochrane reviews) found during the selection process to perform backward and forward citation searching to ensure rigorous coverage of the literature.

Study selection

We used the systematic review management software Distiller-SR (Evidence Partners, Ottawa, Canada) to facilitate study selection. We conducted calibration training of the screening for the first 100 references identified in the literature search to calibrate eligibility criteria interpretation between reviewers. If inter-rater agreements (kappa statistic) were below $\mathbf{k}=0.60$, we clarified the eligibility criteria and conducted a new calibration training on 100 references. Pairs of reviewers independently (FN, JD, MD, OD, RB, TS) screened titles and abstracts of all potentially eligible citations. Subsequently, pairs of

Table 1 Study eligibility criteria.

Category	Study Eligibility Criteria
Population	Adults, adolescents, and children living with RMDs or their important others (e.g., family member, caregiver, friend) involved in the SDM process.
Concept	All measurement instruments used in SDM studies (where SDM was a primary or secondary objective) to assess any SDM outcomes.
Context	Any clinical and research contexts used in SDM studies.
Study designs	Inclusions:
	 quantitative and qualitative primary studies of any designs
	 validation studies of measurement properties
	Exclusions:
	literature reviews
	 protocols
	 conference abstracts, oral presentations
	 editorials, letters, and commentaries
	 studies of any designs available as a thesis
	 consensus statements
Language of publication	Any languages.

reviewers then independently screened the full text of potential studies. Disagreement between pairs of reviewers were resolved by reaching a consensus through discussion within the review team. We performed handsearching of the reference lists of the included studies and the literature reviews found during the selection process to screen for relevant studies not found with our search strategy. We documented reasons for full text exclusion and reported them using PRISMA 2020 flowchart.

Data extraction

We developed a charting form in Distiller-SR to extract data from the articles identified. Before charting began, reviewers made calibration by testing independently the charting form on a random sample of 10 included studies to ensure a mutual understanding of the variables to extract and that the form adequately captured the desired information. Pairs of reviewers performed the data extraction. We resolved disagreements by team consensus. Extracted variables are presented in Table 2.

Data charting and synthesis

Study characteristics and methodology

Objective #1:

We conducted a descriptive analysis centred on the characteristics of

Extracted variables

Year of publication

· Study design

• RMD diagnosis

Sample size

Ethnicity

· Country of publication

· Gender or sex according to the reporting

Objective #1: To identify all available measurement instruments

Table 2
Extracted variables.

Category

Population

Concept	Presence of any variables known to impact patients' involvement in decision making (e.g., income, emotional distress) [21,22] Any measurement instruments used for data collection of SDM outcomes
Objective #2: To explore evidence ga	nps
Category	Extracted variables
Information reported in the included studies that could be useful to explore the three pillars of the OMERACT Filter 2.2.	TRUTH Domain match: Reported measured constructs and their reported definition [12] Reported subscales [12] Construct validity: Any reported information on construct validity [12] DISCRIMINATION Any reported information on: Test-retest reliability or internal consistency [23]. Longitudinal construct validity [23]. Clinical trial discrimination [23]. Thresholds of meaning (e.g., Minimal Important Difference, benchmarks of meaningful scores) [23]. FEASIBILITY Reported number of items [12] Reported scoring and cut-off [12] Type of measure [12] Other reported information on feasibility [12,24]

the included studies and the SDM measurement instruments used in each study.

Objective #2:

We classified the identified instruments according to the presence or absence of any measurement properties reported in the included studies. For instruments without information on measurement properties, we described them to explore evidence gaps for domain matching (i.e., one step of the truth pillar) due to the absence of any information for the two remaining pillars.

For instruments with any measurement properties, we described them to explore evidence gaps for the three pillars of the OMERACT Filter 2.2 (i.e., truth, discrimination, and feasibility). We also conducted an exploratory analysis of evidence gaps related to measurement properties (i.e., truth and discrimination pillars of the OMERACT Filter 2.2) based on an adapted version of the Summary of Measurement Properties (SOMP) table. The adapted SOMP table relies on 7-criteria: feasibility [12], construct validity [12], inter-method reliability [12], test-retest reliability [12], longitudinal construct validity [12], clinical trial discrimination [12], and thresholds of meaning [12].

Results

Study selection

Fig. 1 presents a flow diagram of the selection process (PRISMA flowchart). We retrieved a total of 14,464 citations from our search. Six additional articles were included from hand searching. After removing duplicates, 9905 citations remained for the selection. Of these citations, 239 were retained for full text screening. From these, we found 99 eligible studies that were included for data extraction.

Characteristics of the included studies

Appendix A describes the characteristics of the 99 included studies [25–123]. Fifty-two were performed in the United States, 11 in Canada, and 9 in the Netherlands. Most studies used quantitative designs such as clinical trials or cohort studies (n=80). The five most common RMDs were knee osteoarthritis (OA) (n=36), hip OA (n=20), rheumatoid arthritis (n=18), osteoporosis (n=9), and lumbar herniated disc (n=7). The sample size per study ranged from 11 to 5751 participants. Three

studies only recruited children. From the 96 studies on adult participants, 93 reported information on age (mean age ranged from 24 to 77 years) and sex (female percentage ranged from 5% to 100%), and 49 provided information on ethnicity. From the three studies on children, two reported mean age (ranged from 12 to 13 years with minimum=6 and maximum=17 years), only one study reported information on sex (57% of female). No study presented data on ethnicity or other individual characteristics associated with SDM. The five most reported individual characteristics were information on educational level (n=75), employment status (n=35), marital status (n=29), income (n=20), and access to health insurance (n=15).

We identified 220 candidate measurement instruments associated with SDM outcomes (Appendix A). The five most used measurement instruments were the Decisional Conflict Scale (traditional and low literacy versions) (n=38), the Hip/Knee-Decision Quality Instrument (n=20), the Decision Regret Scale (n=9), the Preparation for Decision Making Scale (n=8), and the Collaborate (n=8). The 80 quantitative studies (i.e., study designs likely to use a Core Outcome Measurement Set) used on average 3.5 (SD=1.9) different SDM measurement instruments

Characteristics of the measurement instruments with any measurement properties reported by the included studies

We identified 44 out of 220 measurement instruments (20%) for which included studies reported any measurement properties (Appendix B). We extracted all reported constructs for each of the instruments. Based on the available data, 15 measurement instruments (34%) reported multiple and conflicting constructs for a given instrument (i.e., different definitions for the same instrument). For example, the Decisional Conflict Scale included studies reporting different constructs such as "decisional conflict", "decision quality" or "aspect of the decision making process".

The extracted instruments were mainly patient-reported outcome measures (n=43, 98%). The number of items per measurement instrument ranged from 1 to 60. Nine studies (21%) reported conflicting information about the number of items and/or subscales. Eight studies (18%) reported conflicting information about the score or cut-off. Thirty-eight (86%) measurement instruments reported information on reliability, 30 (68%) on validity, 6 (14%) on longitudinal construct

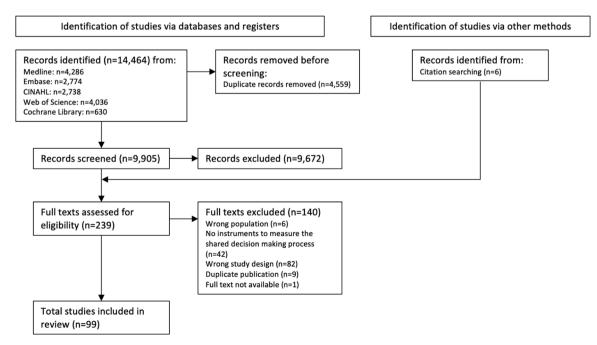


Fig. 1. Completed PRISMA Flowchart.

validity, 6 (14%) on feasibility, and 5 (11%) on thresholds of meaning. From the 99 included studies, only 19 (19%) yielded the populations in which one or more reported measurement instruments were validated. From the data reported by the included studies, only 18 (41%) measurement instruments were partially tested for validation in a RMD sample.

Characteristics of the measurement instruments without any measurement properties reported by the included studies

We identified 176 measurement instruments (80%) for which included studies did not report any measurement properties (Appendix C). We organized these measurement instruments into clusters representing 13 themes. The five largest themes involved

Table 3Exploratory analysis of evidence gaps from an overall summary based on the adapted SOMP.

	Measuremen	nt properties						
Measurement instrument	Feasibility	Truth		Discrimination				
		Construct validity	Inter-method reliability	Test-retest reliability	Longitudinal construct validity	Clinical trial discrimi-nation	Thresholds of meaning	
Decisional Conflict Scale			N/A	X	X	Х	Х	
Decisional Conflict Scale (low literacy)		X	N/A	X		X	X	
SURE Test		X*	N/A			X*	X	
Preparation for Decision Making Scale		X	N/A			X		
Knee-Decision Quality Instrument	X*	X*	N/A	X*	X	X*	X*	
Hip-Decision Quality Instrument	X*	X*	N/A	X*	X	X*	X*	
Herniated disc-Decision Quality Instrument	X*	X*	N/A	X*	X	X*		
Spinal Stenosis-Decision Quality Instrument	X		N/A	X*	X			
ReproKnow	X*	X*	N/A					
Methotrexate in rheumatoid arthritis knowledge test			N/A	X*				
Osteoporosis patient knowledge			N/A					
questionnaire Pregnancy in rheumatoid arthritis			N/A					
questionnaire			14/11					
CollaboRATE		X*	N/A	X (intra-rater)	X	X		
Control Preference Scale		X	N/A N/A	A (IIIIIa=IaiCI)	41	Λ		
Trust in Physician Scale		Α	N/A					
interpersonal Processes of Care			N/A					
Medication adherence			N/A					
Satisfaction With Decision Scale		X	N/A					
Questionnaire on Doctor-Patient Interaction (FAPI)	X*	А	N/A			X *		
9-item Shared Decision Making Questionnaire		X	N/A					
Princess Margaret Hospital Satisfaction with Doctor Questionnaire			N/A					
nformed Shared Decision Making Scale			N/A					
Decision Regret Scale		X	N/A					
Shared Decision Making Process		X*	N/A	X		X*		
rust in Surgical Decision Scale			N/A					
Decision Self Efficacy Scale		X	N/A			X		
Patient-Doctor Relationship Questionnaire			N/A					
OPTION Scale								
MASRI		X	N/A					
Beliefs about Medicines Questionnaire		X	N/A	X				
nterpersonal Trust in a Physician			N/A	X				
Effective Consumer Scale			N/A	X				
Medication Education Impact Ouestionnaire			N/A	X*				
Morisky Medication Adherence Scale			N/A					
Perceived Involvement in Care Scale			N/A					
Satisfaction with Information about Medicines Scale			N/A					
Decision Evaluation Scales			N/A					
Cologne Patient Questionnaire			N/A					
Decision readiness			N/A					
Stage of Decision Making Scale			N/A					
Treatment intention			N/A	X	X			
Satisfaction with decision and decision making			N/A					
Decision process			N/A	X*				
Knowledge on acute low back pain			N/A					

N/A: non-applicable

FAPI: Fragebogen zur Arzt-Patienten-Interaktion

OPTION Scale: Observing Patient Involvement in Decision Making instrument

MASRI: Medication Adherence Self-Report Inventory

^{*} amongst people with RMDs

instruments reporting the measure of satisfaction (n=31, 18%), decision (n=23, 13%), values and preferences (n=22, 13%), adherence (n=22, 13%), and knowledge (n=20, 11%). For the construct of knowledge (the first OMERACT Core Domain [16]), 13 measurement instruments (65%) were specific to one decision in one condition (e.g., a questionnaire about acupuncture in low back pain, a questionnaire about rheumatoid arthritis medications, a questionnaire about total knee replacement). Measurement instruments were mainly patient-reported outcome measures (n=105, 60%). Only 11 measurement instruments (6%) were presented with a name (e.g., Choice Predisposition Scale, Partners in Health Scale) hinting that the remaining instruments were homemade. The number of items per measurement instrument ranged from 1 to 27.

Exploratory analysis of evidence gaps from an overall summary based on the adapted SOMP

Table 3 presents an exploratory analysis of evidence gaps from an overall summary based on the adapted SOMP. We determined matching between the included studies reporting any measurement properties and any of the 7-criteria of the adapted SOMP. From the 44 candidate instruments with any reported measurement properties, 25 (57%) reported at least one measurement property required for the adapted SOMP. Feasibility was reported by 6 out of 25 (24%) instruments, one or more hypotheses testing for construct validity by 16 (64%) instruments, test-retest reliability by 15 (60%) instruments, longitudinal construct validity by 6 (24%) instruments, clinical trial discrimination by 11 (44%) instruments, and thresholds of meaning by 5 (20%) instruments.

From the available data, only 12 out of 44 (27%) instruments reported validation process in RMDs populations. From these 12 instruments, feasibility on RMDs populations was reported by 5 (42%) instruments, one or more hypotheses testing for construct validity by 7 (58%) instruments, test-retest reliability by 7 (58%) instruments, longitudinal construct validity by no instrument, clinical trial discrimination by 6 (50%) instruments, and thresholds of meaning by 2 (17%) instruments. From the 99 included studies, twenty-four (24%) studies only reported that the "instrument is validated", which was not considered sufficient to match with any adapted SOMP criteria.

Discussion

In this scoping review, we were able to identify 220 candidate instruments for SDM outcomes in RMDs. Our classification of the instruments identified evidence gaps prior to conducting the OMERACT Filter 2.2. This led us to make observations concerning 1- the usefulness of scoping review methods to identify candidate instruments in the development of a Core Outcome Measurement Set and 2- inconsistent reporting of key metrics to develop a Core Outcome Measurement Set under a high-quality framework such as the OMERACT Filter 2.2.

First, to our knowledge, this is the first scoping review to identify the broad diversity of SDM instruments used in the field of RMDs. We are amongst the first OMERACT Working Group to use a scoping review to identify candidate instruments to develop a Core Outcome Measurement Set within the OMERACT Filter 2.2 framework. We hypothesized that the two specific objectives of scoping review (i.e., data charting and research evidence gaps) could be of great value in informing the operationalization of the OMERACT Filter 2.2 in the field of SDM [18,124]. This scoping review did not seek to provide definitive answers to the OMERACT Filter 2.2, but rather is an opportunity to explore key variables reported by authors and evidence gaps that could require further investigation prior to the OMERACT Filter 2.2.

We identified over 200 candidate instruments, demonstrating the scale and complexity of measuring SDM. This high number of candidate instruments contrasts with the Working Group's aim of selecting only a few standardized tools with high measurement properties value to be used across all SDM trials in RMDs. Our results showed that clinical studies within our sample of eligible articles only used on average four

SDM measurement instruments. We also identified clusters of instruments used in more studies such as the Decisional Conflict Scale, a recognized standard in the field of SDM [3], or the Hip/Knee-Decision Quality Instrument [100–103,105,106]. However, these instruments do not appear to cover all of six OMERACT Core Domains [16] and we are uncertain about the full scale of their relevant measurement properties. Also, the number of items varied greatly between instruments. These findings will need to be accounted for in the selection process of the instruments using the OMERACT Filter 2.2.

Our scoping review will serve as a repository of possible measures of SDM in RMDs studies. This is an opportunity not to bias the selection of the Core Outcome Measurement Set on a few "legacy measures" in the field of SDM, but rather give full opportunity to multiple stakeholders to express their preferences concerning the measure of SDM in RMDs. The SDM Working Group will need to assess the relevance of all candidate instruments and triage them according to the consensus-based Core Domains. A limited number of instruments will need to be selected to ensure the feasibility and acceptability of the future Core Outcome Measurement Set, as evidence showed that uptake is still limited in clinical trials even for established Core Outcome Measurement Sets [125–127].

Second, we identified evidence gaps and inconsistent reporting of key metrics for the development of a Core Outcome Measurement Set such as measured constructs and measurement properties. Measurement properties of candidate instruments are a vital metric to assess the value of an instrument in the OMERACT Filter 2.2. Unfortunately, we found that only a fifth of all candidate instruments had any measurement properties reported in the included studies. It is possible that this finding exposes inconsistency or underreporting of evidence supporting the use of an instrument in a clinical study. However, the extent of this underreporting highlights that in the SDM literature for RMDs, most authors included multiple measurement instruments without evidence of measurement properties. In the case of our working group and with the current data, we are unable to complete the second step of the OMERACT Filter 2.2 with over 80% of the identified candidate instruments because of the lack of measurement properties.

Another worrying result is that a third of all instruments with any measurement properties reported multiple and conflicting constructs for a given instrument. For example, we extracted six different constructs for our most used instrument the Decisional Conflict Scale, such as "Decisional conflict", "Aspect of the decision making process" or "Perception of being uncertain, uninformed, unsupported, or unclear as to values to be considered". For the Hip/Knee-Decision Quality Instrument, the second most used instrument, we extracted ten different constructs. We also found inconsistent reporting of the number of items and/or subscales used for an instrument and variation in scores and cutoff value interpretation with or without supporting evidence.

Our working group piloted a Core Domain matching exercise at the OMERACT 2023 Special Interest Group meeting. This exercise revealed that several items of the Decisional Conflict Scale were not understood the same way across participants. Domain matching is also compromised for the domain "Knowledge" (i.e., first OMERACT Core Domain). The 20 candidate instruments found in our scoping review are condition-specific and/or decision-specific. Developing a Core Outcome Measurement Set with a unique generic measurement instrument to assess the knowledge component could be difficult and irresponsive to the researchers' needs. Our working group must clarify whether we want a unique instrument for this domain. Given the disparities in reporting constructs for all candidate instruments, we will require a specific methodology to decide, with experts in SDM, which instrument match which Core Domains and which are feasible to complete the first step of the OMERACT Filter 2.2.

To prepare the OMERACT Filter 2.2, we completed an adapted SOMP to verify if the available measurement properties were sufficient to match with any of the 7-criteria of the SOMP as would be required for an instrument to be considered valid for inclusion in an OMERACT Core

Outcome Measurement Set. Again, only half of the instruments with any measurement properties reported at least one measurement property required for the adapted SOMP. This situation may be due to lack of reporting or unavailability of the measurement property for an instrument but demonstrate another significant gap to solve. An example was the Decisional Conflict Scale that could not be considered "Feasible" based on currently available data from our included studies, while other instruments had this information. We are aware of two scoping reviews only for the Decisional Conflict Scale which was used in over 200 studies [128,129] outside the field of RMDs. Our operational definition of this adapted SOMP criteria might be too stringent for SDM instruments [12, 24], but this illustrates evidence gaps for future steps of the process.

Overall, evidence gaps in the reporting of key metrics currently limit the SDM OMERACT Working Group to fully complete the OMERACT Filter 2.2 with certainty. We thereby propose the following possible solutions prior to conducting the Filter 2.2. First, the group will require a consensus-based methodology to match instruments with Core Domains and measure feasibility with potential users. With an organized list of instruments, the group will then require systematic reviews of the identified instrument to ensure all validation studies, including primary development studies, are identified within or outside the field of RMDs. It is possible that some instruments will have been thoroughly validated in other conditions and would be readily available to guide the Filter 2.2. If reviews fail to identify measurement properties required by the SOMP table, we could validate some of these instruments in RMDs population using our international group's databases, trials, and cohorts in rheumatology. It is also possible that we will require to develop a new instrument that will be feasible to assess all Core Domains. While this may appear a tenuous process that could delay the availability of a Core Outcome Measurement Set for SDM trials, experts in SDM will be unsurprised by the complexity of measuring six different Core Domains, and this work will have reaching consequences for this field as no SDM Core Outcome Measurement Set has been designed for any disease yet.

Strengths and limitations

This scoping review used the highest methodological standards to map a diverse and complex literature such as SDM for RMDs. Our search strategy allowed the identification of candidate instruments used in various SDM interventions, for multiple RMDs and using a wide spectrum of study designs. However, the complexity of the key concept "SDM" and its heterogeneity may have hindered identification of some studies on this topic. The lack of standardized search filter for RMDs may have also missed relevant studies due to the wide range of diagnostic labels. Our information specialist had previously designed search strategies in other high-quality reviews on SDM and we are confident that we identified most relevant studies and possible instruments to consider for developing a future Core Outcome Measurement Set within the OMERACT framework.

The main limitation is that the scoping review does not enable the SDM Working Group to fully complete the OMERACT Filter 2.2 in its current form. Since completing the Filter 2.2 requires an in-depth analysis of measurement properties, this information is only obtainable if the authors correctly reported any measurement properties in the included studies. The scoping methodology is not able to identify all validation studies for every instrument. We suspect that many instruments considered "unvalidated" in this review, may be so from a lack of complete reporting of measurement properties in clinical studies having used them. Systematic reviews outside the field of RMDs may be required to fully describe measurement properties of selected candidate instruments.

Conclusion

We identified 220 candidate instruments used in the SDM literature amongst people with RMDs. Our classification of instruments showed

evidence gaps and inconsistent reporting of measured constructs and measurement properties. The evidence gaps currently limit the capacity to fully complete the OMERACT Filter 2.2 with certainty. The next steps for the OMERACT SDM Working Group are to match candidate instruments with Core Domains, assess feasibility and review validation studies of measurement instruments in rheumatic diseases or other conditions. Development and validation of new instruments may be required for some Core Domains.

Funding

This study was funded by the Canadian Institutes of Health Research.

CRediT authorship contribution statement

Florian Naye: Conceptualization, Methodology, Investigation, Writing - original draft. Karine Toupin-April: Conceptualization, Methodology, Investigation, Writing - original draft. Maarten de Wit: Investigation, Writing - review & editing. Annie LeBlanc: Conceptualization, Methodology, Investigation, Writing - original draft. Olivia Dubois: Conceptualization, Methodology, Investigation, Writing original draft. Annelies Boonen: Investigation, Writing - review & editing. Jennifer L. Barton: Investigation, Writing – review & editing. Liana Fraenkel: Investigation, Writing – review & editing. Linda C. Li: Investigation, Writing – review & editing. Dawn Stacey: Investigation, Writing - review & editing. Lyn March: Investigation, Writing - review & editing. Claire E.H. Barber: Investigation, Writing – review & editing. Glen Stewart Hazlewood: Investigation, Writing - review & editing. Francis Guillemin: Investigation, Writing - review & editing. Susan J. Bartlett: Investigation, Writing - review & editing. Dorthe B. Berthelsen: Investigation, Writing - review & editing. Kate Mather: Investigation, Writing - review & editing. Laurent Arnaud: Investigation, Writing - review & editing. Akpabio Akpabio: Investigation, Writing - review & editing. Adewale Adebajo: Investigation, Writing review & editing. Grayson Schultz: Investigation, Writing - review & editing. Victor S. Sloan: Investigation, Writing - review & editing. Tiffany K. Gill: Investigation, Writing - review & editing. Saurab Sharma: Investigation, Writing - review & editing. Marieke Scholte-Voshaar: Investigation, Writing - review & editing. Francesco Caso: Investigation, Writing - review & editing. Elena Nikiphorou: Investigation, Writing - review & editing. Samah Ismail Nasef: Investigation, Writing - review & editing. Willemina Campbell: Investigation, Writing - review & editing. Alexa Meara: Investigation, Writing - review & editing. Robin Christensen: Investigation, Writing – review & editing. Maria E. Suarez-Almazor: Investigation, Writing - review & editing. Janet Elizabeth Jull: Investigation, Writing – review & editing. Rieke Alten: Investigation, Writing – review & editing. Esi M. Morgan: Investigation, Writing - review & editing. Yasser El-Miedany: Investigation, Writing - review & editing. Jasvinder A. Singh: Investigation, Writing - review & editing. Jennifer Burt: Investigation, Writing - review & editing. Arundathi Jayatilleke: Investigation, Writing – review & editing. Ihsane Hmamouchi: Investigation, Writing - review & editing. Francisco J. Blanco: Investigation, Writing - review & editing. Anthony P. Fernandez: Investigation, Writing - review & editing. Sarah Mackie: Investigation, Writing - review & editing. Allyson Jones: Investigation, Writing - review & editing. Vibeke Strand: Investigation, Writing - review & editing. Sara Monti: Investigation, Writing - review & editing. Simon R. Stones: Investigation, Writing review & editing. Rebecca R. Lee: Investigation, Writing - review & editing. Sabrina Mai Nielsen: Investigation, Writing – review & editing. Vicki Evans: Investigation, Writing - review & editing. Hemalatha Srinivasalu: Investigation, Writing - review & editing. Thomas Gérard: Investigation, Writing - review & editing. Juliette LeBlanc Demers: Conceptualization, Methodology, Investigation, Writing original draft. Roxanne Bouchard: Conceptualization, Methodology, Investigation, Writing – original draft. Théo Stefan: Conceptualization,

Methodology, Investigation, Writing – original draft. Michèle Dugas: Conceptualization, Methodology, Investigation, Writing – original draft. Frédéric Bergeron: Methodology. Dorcas Beaton: Investigation, Writing – review & editing. Lara J. Maxwell: Conceptualization, Methodology, Investigation, Writing – review & editing. Peter Tugwell: Conceptualization, Methodology, Investigation, Writing – review & editing. Simon Décary: Conceptualization, Methodology, Investigation, Writing – original draft.

Declaration of competing interest

The authors declare the following financial interests/personal relationships which may be considered as potential competing interests:

Anthony P. Fernandez, MD, PhD: Past 36 months: Grants or contracts from any entity: Mallinckrodt, Novartis, Pfizer. Payments to institution and (partial) to me. Consulting fees: AbbVie, Biogen, UCB, BMS, Alexion: Payments to me. Payment or honoraria for lectures, presentations, speakers bureaus, manuscript writing or educational events: AbbVie, BMS, Kyowa Kirin, Mallinckrodt: Payments to me. Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid: Board of Directors, American Society of Dermatopathology; Associate Editor, Journal of the American Academy of Dermatology.

Arundathi Jayatilleke: Pas 36 months: Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid: Pennsylvania Rheumatology Society, board member: unpaid.

Claire Barber: Pas 36 months: Grants or contracts from any entity: CIHR – 3: Peer reviewed national funding unrelated to current project. CIORA (Canadian Rheumatology Association): Peer reviewed national funding unrelated to current project. Cumming school of medicine Seed Grant: Local university funding, peer reviewed, unrelated to current project. Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid: Past chair Human Resource Committee, Canadian Rheumatology Association: Unrelated to current project.

Dorcas Beaton: Past 36 months: Support for attending meetings and/or travel: OMERACT Management Team: OMERACT covers travel costs for members of management team to attend conferences on behalf of OMERACT. Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid: Member of Management team of OMERACT, chair of methodology at OMERACT: I help make decisions on the methods that will be used to come to a decision about core outcome sets at OMERACT. This would have informed methods used in this paper.

Dorthe B Berthelsen: Past 36 months: Grants or contracts from any entity: Have received PhD Scholarships from Odense University Hospital and from the Faculty of Health Sciences, University of Southern Denmark. Support for attending meetings and/or travel: Have received a grant from the Erna Hamilton Foundation to cover meeting registration fee and travel costs for OMERACT 2023.

Dawn Stacey: Past 36 months: Grants or contracts from any entity: Canadian Institutes of Health Research. Support for attending meetings and/or travel: Beijing University of Chinese Medicine – August 2023. Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid: Co-Chair International Patient Decision Aid Standards Collaboration (unpaid).

Karine Toupin-April: Past 36 months: Support for attending meetings and/or travel: OMERACT travel award given to the Shared decision making group to help attend the OMERACT 2023 meeting.

Lyn MARCH: Past 36 months: Grants or contracts from any entity: Commonwealth Government of Australia Medical Research Future Fund: RCT for biological tapering in RA and PsA (Utilising shared decision making): Payment to institution. Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid: Chair, Australian Rheumatology Association Research Fund Committee: unpaid. Executive, Global Alliance for MSK Health: unpaid.

Maria Suarez-Almazor: Past 36 months: Consulting fees: Pfizer:

Consultant. Eli Lilly: Consultant. Syneos Health: Consultant. Participation on a Data Safety Monitoring Board or Advisory Board: Celgene: DSMB member.

Maarten de Wit: Past 36 months: Consulting fees: UCB: Payment to Stichting Tools, Netherlands.

Peter Tugwell: Past 36 months: Consulting fees: Reformulary Group: Providing independent medical consultation professional services to the firms listed in this section. Participation on a Data Safety Monitoring Board or Advisory Board: UCB Biopharma GmbH & SPRL, Parexel International, Prahealth Sciences: An independent Committee Member for clinical trial Data Safety Monitoring Boards for FDA approved trials being conducted by: - UCB Biopharma GmbH & SPRL, - Parexel International, - Prahealth Sciences. Other financial or non-financial interests: Abbvie, Astra Zeneca, Aurinia, BMS, Centrexion, GSK, Horizon Pharma Inc, Janssen, Novartis, Pfizer & Sparrow: I am [unpaid] Chair of the Management Group of a registered non-profit independent medical research organization, OMERACT, whose goal is to improve and advance the health outcomes for patients suffering from musculoskeletal conditions. OMERACT receives arms-length funding from 11 companies.

Rieke Alten: Past 36 months: Consulting fees: Abbvie, BMS, CELLT-RION; Eli Lilly; Galapagos, Janssen, Lilly, Novartis, Pfizer, Roche, UCB, Viatris. Payment or honoraria for lectures, presentations, speakers bureaus, manuscript writing or educational events: Abbvie, BMS, CELLT-RION; Eli Lilly; Galapagos, Janssen, Lilly, Novartis, Pfizer, Roche, UCB, Viatris. Support for attending meetings and/or travel: Abbvie, BMS, CELLTRION; Eli Lilly; Galapagos, Janssen, Lilly, Novartis, Pfizer, Roche, UCB, Viatris. Participation on a Data Safety Monitoring Board or Advisory Board: Abbvie, BMS, CELLTRION; Eli Lilly; Galapagos, Janssen, Lilly, Novartis, Pfizer, Roche, UCB, Viatris.

Susan J. Bartlett: Past 36 months: Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid: American Thoracic Society Board of Directors: 2020–2022; unpaid. PROMIS Health Organization – President Elect, Board of Directors: unpaid. American College of Rheumatology Association of Rheumatology Professionals Executive Committee: 2021–2023; unpaid.

Simon Stones: Past 36 months: Consulting fees: Future Science Group: Payment for document review. Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid: RAiISE: Director. Stock or stock options: Envision Pharma Group: Related to employment. Other financial or non-financial interests: Envision Pharma Group: Employment.

Esi Morgan: Past 36 months: Grants or contracts from any entity: Pfizer, Inc: Educational Program to Optimize Delivery of Care to Families with Juvenile Idiopathic Arthritis Over Telemedicine; Investigator Initiated Grant to Seattle Children's Research Institute, role - co-Investigator. Agency for Healthcare Research and Quality: Informing personalized treatment decision with advanced Bayesian causal inference - A patient-centred evidence-based shared decision making (SDM) digital health technology; Investigator Initiated Grant to Seattle Children's Research Institute, role - PI (multi-PI grant). Payment or honoraria for lectures, presentations, speakers bureaus, manuscript writing or educational events: American College of Rheumatology: Honorarium, Education Conference April 2023. Support for attending meetings and/ or travel: American College of Rheumatology: Travel Support Education Exchange Conference April 2023. Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid: pediatric Rheumatology Care and Outcomes Improvement Network: Principal Investigator.

Francis GUILLEMIN: Past 36 months: Grants or contracts from any entity: Novartis: Payment to my institution.

Hemalatha Srinivasalu: Past 36 months: Grants or contracts from any entity: CARRA Registry Associate and NIAMS Intramural program.

Janet Jull: Past 36 months: Grants or contracts from any entity: Canadian Institutes of Health Research.

Jennifer L. Barton: Past 36 months: Grants or contracts from any entity: US Department of Veterans Affairs.

Jasvinder A. Singh: Past 36 months: Consulting fees: Crealta/Horizon, Medisys, Fidia, PK Med, Two labs Inc., Adept Field Solutions, Clinical Care options, Clearview healthcare partners, Putnam associates, Focus forward, Navigant consulting, Spherix, MedIQ, Jupiter Life Science, UBM LLC, Trio Health, Medscape, WebMD, and Practice Point communications; and the National Institutes of Health and the American College of Rheumatology: Consultant fees paid to me for each entity. Payment or honoraria for lectures, presentations, speakers bureaus, manuscript writing or educational events: JAS is on the speaker's bureau of Simply Speaking: Consultant fees paid to me. Support for attending meetings and/or travel: Past steering committee member of OMERACT: I previously received support from the organization to attend their meeting every 2 years. Participation on a Data Safety Monitoring Board or Advisory Board: FDA Arthritis Advisory Committee: JAS serves as a member. No financial support. Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid: Past steering committee member of the OMERACT, an international organization that develops measures for clinical trials and receives arms length funding from 12 pharmaceutical companies: I previously received support from the organization to attend their meeting every 2 years. Co-Chair of the Veterans Affairs Rheumatology Field Advisory Committee: No financial support. Editor and the Director of the UAB Cochrane Musculoskeletal Group Satellite center on Network Meta-analysis: No financial support. Stock or stock options: JAS owns stock options in Atai life sciences, Kintara therapeutics, Intelligent Biosolutions, Acumen pharmaceutical, TPT Global Tech, Vaxart pharmaceuticals, Atyu biopharma, Adaptimmune Therapeutics, GeoVax Labs, Pieris Pharmaceuticals, Enzolytics Inc., Seres Therapeutics, Tonix Pharmaceuticals Holding Corp., and Charlotte's Web Holdings, Inc.: I own stock options. JAS previously owned stock options in Amarin, Viking and Moderna pharmaceuticals: I owned stock options in these companies previously.

Saurab Sharma: Past 36 months: Grants or contracts from any entity: I am supported by the International Association for the Study of Pain John J. Bonica Postdoctoral Fellowship. The funder does not have any influence on my research. Support for attending meetings and/or travel: My travel was supported to present a talk (unrelated to the manuscript) at the International Association for the Study of Pain (IASP) Congress in Toronto in 2022.

Victor Sloan: Past 36 months: Consulting fees: Boehringer-Ingelheim: Payment to me. Stock or stock options: Stock in UCB Pharma.

Willemina Campbell: Past 36 months: Payment or honoraria for lectures, presentations, speakers' bureaus, manuscript writing or educational events: ABBVIE and Einstein Medical School. Support for attending meetings and/or travel: OMERACT and GRAPPA. Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid: GRAPPA PRP CHAIR-past.

Acknowledgments

We would like to recognize the valuable contribution of the OMERACT Executive team Shawna Crosskleg and Bev Shea. We take Marianne Ruel for conducting the PRESS checklist.

Supplementary materials

Supplementary material associated with this article can be found, in the online version, at doi:10.1016/j.semarthrit.2023.152344.

Appendix A. Characteristics of the 99 included studies

Authors Year of publica- tion	Country	Study design	Condition(s)	Sam- ple size (n)	Characteristics of participants Age=mean (SD) [Range]	SDM (instrument) measures
Allen et al. 2016 [25]	United States of America	Quantitative	Hip or Knee OA	155	 Age: 61.8 (11.7) Female: 60.6% Ethnicity: 58.1% Caucasian, 38.7% African American, 3.2% Other Information on education level, numeracy, literacy, health insurance 	Decisional Conflict Scale (low literacy) Preparation for Decision Making Scale Knee-Decision Quality Instrument Hip-Decision Quality Instrument Stage of Decision Making Scale
Andersen et al. 2019 [26]	Denmark	Quantitative	Lumbar herniated disc	40	Age: NRFemale: NREthnicity: NR	Decisional Conflict Scale CollaboRATE Herniated Disc-Decision Quality Instrument
Bansback et al. 2022 [27]	Canada	Quantitative	Knee OA	163	 Age: 64.17 (8.34) to 64.95 (7.54) Female: 46.3% to 64.2% Ethnicity: NR Information on emotional distress 	Knee-Decision Quality Instrument SURE Test CollaboRATE Control Preference Scale Value concordance analysis Item on treatment preference Item on willingness to have surgery
Barton et al. 2014 [28]	United States of America	Quantitative	Rheumatoid arthritis	509	 Age: 55 (14) to 64 (11) Female: 84% to 86% Ethnicity: 36–83% White, 8–32% Latino, 5–20% Asian/Pacific Islander, 1–8% African American, 3–4% Other Information on education level, income, literacy, native language, emotional distress 	Trust in Physician Scale Interpersonal Processes of Care (continued on next page)

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Authors Year of publica- tion	Country	Study design	Condition(s)	Sam- ple size (n)	Characteristics of participants Age=mean (SD) [Range]	SDM (instrument) measures
Barton et al. 2016 [29]	United States of America	Quantitative	Rheumatoid arthritis	166	 Age: 58 (12) [24–85] Female: 88% Ethnicity: 45% Latino, 26% Asian, 14% African American, 13% Caucasian, 3% Other Information on marital status, employment, emotional distress 	Decisional Conflict Scale (Low literacy) Trust in Physician Scale Interpersonal Processes of Care Item on medication adherence Knowledge questionnaire about rheumatoid arthritis medications
Bieber et al. 2008 [30]	Germany	Quantitative	Fibromyalgia	85	 Age: 49.5 (11.3) to 50.4 (8.8) Female: 90.2% to 93.2% Ethnicity: NR Information on education level, income, literacy, native language, emotional distress 	 Decisional Conflict Scale Satisfaction With Decision scale Questionnaire on Doctor- Patient Interaction (FAPI)
Birru Talabi et al. 2019 [31]	United States of America	Validation	Rheumatic diseases	152	Age: [18–50] Female: 100% Ethnicity: 77.1% White, 22.2% Non-white Information on education level	• ReproKnow
Bishop et al. 2019 [32]	United Kingdom	Quantitative	Back pain	350	 Age: 47.9 (15.8) Female: 56.3% Ethnicity: 88.9% White British, 4.6% White other, 1.2% Asian or Asian British, 0.6 Mixed, 0.6 Black or Black British Information on education level 	Knowledge questionnaire about acupuncture Item on willingness to have acupuncture
Boland et al. 2018 [33]	Canada	Quantitative	Knee OA	242	Age: 65 (10.3) to 69 (8.2) Female: 51% to 63% Ethnicity: NR Information on education level, employment, income, native language	Knee-Decision Quality Instrument SURE Test
Bossen et al. 2022 [34]	Netherlands	Quantitative	Hip or knee OA	317	 Age: 68 (8.69) to 71 (8.28) [46–90] Female: 49.71% to 51.7% Ethnicity: NR Information on education level 	 Decisional Conflict Scale Patient Activation Measure 9-item Shared Decision- Making Questionnaire
Bot et al. 2014 [35]	United States of America	Mixed methods	Nontraumatic painful conditions of the upper extremity	130	 Age: 52 (16) [18–91] Female: 52% Ethnicity: NR Information on marital status, education level, employment 	 Princess Margaret Hospital Satisfaction With Doctor Questionnaire Informed Shared Decision Making scale
Bozic et al. 2013 [36]	United States of America	Quantitative	Hip or knee OA	123	Age: 63.1 (10.5) [19-85] Female: 54.5% Ethnicity: 73.2% White, 8.1% Asian, 4.9% Black or African American, 0.8% American Indian or Alaska native, 0% Native Hawaiian or pacific islander, 4.9% other Information on education level, employment, income, health insurance	Knowledge questionnaire about OA of the hip and knee Item on treatment choice Length of consultation time Item on satisfaction with the visit
Braddock et al. 2008 [37]	Canada	Qualitative	Orthopaedic surgery	133	 Age: 71.2 [60–96] Female: 74% Ethnicity: 76% White, 21% Black, 4% Hispanic Information on education level 	Observation tool for informed decision making Duration of the visit
Brinkman et al. 2017 [38]	United States of America	Quantitative	Juvenile idiopathic arthritis (parents reported the information)	171	Age: NRFemale: NREthnicity: NR	SURE testCollaborate
Brodney et al. 2019 [39]	United States of America	Validation	Hip or knee OA, lumbar herniated disc or lumbar spinal stenosis	649	 Age: 59.9 (15.2) to 64.8 (10.8) Female: 42% to 56% Ethnicity: 87–93% White, 2–5% Black, 1–2% Hispanic, 4–6% Other Information on education level 	Decision Regret Scale CollaborATE Shared Decision Making Process SURE Test Item on overall satisfaction Informed choice analysis
Brodney et al. 2022 [40]	United States of America	Quantitative	Hip or knee OA, lumbar herniated disc or lumbar spinal stenosis	700	 Age: 65.8 Female: 55.3% Ethnicity: 89.4% White Information on education level, literacy 	 Trust in the Surgical Decision Scale Decision Regret Scale Shared Decision Making Process Scale (continued on next page)

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Authors Year of publica- tion	Country	Study design	Condition(s)	Sam- ple size	Characteristics of participants Age=mean (SD) [Range]	SDM (instrument) measures
				(n)		
Cranney et al. 2002 [42]	Canada	Quantitative	Osteoporosis	18	 Age: 61.4 (9.74) Female: 100% Ethnicity: NR 	Knee-Decision Quality Instrument Hip-Decision Quality Instrument Herniated Disc-Decision Quality Instrument Spinal Stenosis-Decision Quality Instrument Decisional Conflict Scale Decision Self-Efficacy Scale Knowledge questionnaire
					Information on education level, employment	about osteoporosis and the available treatments Items on realistic expectations Items on values Item on choice predisposition Item on choice
Chen et al. 2021 [41]	Taiwan	Quantitative	Lumbar degenerative diseases	130	 Age: 54.9 to 55.7 Female: 60.3% to 67.2% Ethnicity: NR Information on education level 	 Decisional Conflict Scale Satisfaction With Decision scale 9-item Shared Decision Making Questionnaire Decision Self-Efficacy Scale Control Preference Scale
de Achaval et al. 2012 [43]	United States of America	Quantitative	Knee OA	208	 Age: 62.8 (9.0) Female: 68% Ethnicity: 66% White, 24% African American, 7% Hispanic, 3% Other Information on education level, employment 	Decisional Conflict Scale
de Jesus et al. 2017 [44]	Canada	Quantitative	Knee OA	45	Age: 64.6 [50–90]Female: 42.2%Ethnicity: NR	Decisional Conflict Scale Knowledge questionnaire about knee OA options Item on preparation to make a decision on their preference
Drenkard et al. 2019 [45]	United States of America	Quantitative	Systemic lupus erythemato-sus	698	 Age: 47.5 (13.7) Female: 93.1% Ethnicity: NR Information on education level, employment, health insurance, emotional distress 	Interpersonal Processes of Care
du Long et al. 2016 [46]	Netherlands	Quantitative	Hip or knee OA	172	Age: 65 (11) [31–91]Female: 57%Ethnicity: NR	 Decisional Conflict Scale Patient-Doctor Relationship Questionnaire
El Miedany et al. 2019 [47]	Egypt	Quantitative	Juvenile idiopathic arthritis	189	 Age: 12.7 (1.3) to 12.8 (1.5) [6.1–15.5] Female: 56.8 to 57.4% Ethnicity: NR 	 9-item Shared Decision Making Questionnaire Compliance analysis Persistence analysis
Elwyn et al. 2016 [48]	United Kingdom	Mixed methods	Knee OA	72	 Age: 65.8 (11.3) Female: 60% Ethnicity: NR Information on education level, native language 	 Knee-Decision Quality Instrument OPTION Scale Treatment alignment analysis Item on values
Espinoza et al. 2022 [49]	Canada	Qualitative	Osteoporosis	169	 Age: 57.8 (14.6) Female: 61.5% Ethnicity: 93.4% White/ Caucasian, 3.6% Black/African American, 3% Other Information on marital status, education level, income, health insurance 	 Compliance analysis Persistence analysis
Fraenkel et al. 2007 [52]	United States of America	Quantitative	Pain involving one or both knees	83	 Age: 74 (7) to 74 (9) Female: NR Ethnicity: 65–72% Caucasian, 21–30% African American Information on marital status, education level 	 Decision Self-Efficacy Scale Preparation for Decision Making Tool
Fraenkel et al. 2012 [51]	United States of America	Quantitative	Rheumatoid arthritis	104	 Age: 62 (12) Female: 84% Ethnicity: 87% White Information on education level, employment 	 Decisional Conflict Scale Choice Predisposition Scale Knowledge questionnaire about biologics Informed choice analysis (continued on next page)

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Authors Year of publica- tion	Country	Study design	Condition(s)	Sam- ple size (n)	Characteristics of participants Age=mean (SD) [Range]	SDM (instrument) measures
Fraenkel et al. 2015 [50]	United States of America	Quantitative	Rheumatoid arthritis	121	 Age: 54.3 (11.4) to 56.2 (13.3) Female: 68% to 72% Ethnicity: 97–98% White, 5–7% Hispanic Information on marital status, education level, employment, 	Items on values Decisional Conflict Scale Combined Outcome Measure for Risk Communication Informed choice analysis
Gasteiger et al. 2022 [53]	New Zealand	Quantitative	Rheumatic diseases (rheumatoid arthritis, ankylosing spondylitis, psoriatic arthritis, granulomatosis with polyangiitis, juvenile idiopathic arthritis, other) and their companions	79	literacy • Age: 54.1 (17.1) • Female: 60% • Ethnicity: 61% New Zealand European, 20% Other, 9% Pacific, 8% Asian, 3% Māori • Information on education level	Decisional Conflict Scale Satisfaction With Decision scale Item on willingness to change to a biosimilar Item on preference towards biosimilars Item on perceptions of cognitive and affective risk Items on practical and emotional support received by accompanied patients during the decision process Item on explanation understanding Item on reassurance Item on preferences in receiving information accompanied
Georgo- poulou et al. 2020 [54]	United Kingdom	Quantitative	Lupus nephretis	98	 Age: 40 (10.94) [21–66] Female: 85.7% Ethnicity: 35.7% English, 1% White and Black African, 1% Irish, 4.1% Any other white background, 1% White and Black Caribbean, 2% White and Asian, 2% Any other mixed/multiple background, 3.1% Indian, 7.1% Chinese, 17.3% African, 14.3% Caribbean, 2% any other ethnic background 	Collaborate MASRI (adherence) Beliefs about Medecines Questionnaire Patient-Doctor Relationship Questionnaire Interpersonal Trust in a Physician
Gong et al. 2017 [55]	South Korea	Quantitative	Carpal tunnel syndrome	66	Information on education level Age: 52 (9) to 53 (10) Female: 76% to 81% Ethnicity: NR Information on education level	Decisional Conflict Scale Knowledge questionnaire about carpal tunnel syndrome
Grevnerts et al. 2022 [56]	Sweden	Quantitative	Anterior cruciate ligament injury	101	Age: NR Female: 55% Ethnicity: NR	Items on shared decision making process
Hirata et al. 2023 [57]	Japan	Quantitative	Rheumatic diseases	94	Age: median=66 [52-71]Female: 70%Ethnicity: NR	 Continuance rate of treatment Analysis on influential values of patient
Hochleh-nert et al. 2006 [58]	Germany	Quantitative	Fibromyalgia	75	 Age: 49.85 (10.42) Female: 93.33% Ethnicity: NR Information on marital status, employment 	Decisional Conflict ScaleSatisfaction With Decision scale
Hoffman et al. 2014 [59]	United States of America	Quantitative	Knee OA	126	 Age: [18–85] Female: 61% Ethnicity: 58% Caucasian, 30% African American, 11% Hispanic, 1% Other Information on education level 	Decisional Conflict Scale (low literacy) Preparation for Decision Masking scale Choice Predisposition Scale Knee-Decision Quality Index
Holland et al. 2016 [60]	United States of America	Quantitative	Acute musculoske-letal pain	94	 Age: 70 [60–94] Female: 62% Ethnicity: 74% White, 26% Black Information on education level, literacy 	 Item on satisfaction with decision Item on satisfaction with treatment
Hsiao et al. 2019 [61]	United States of America	Qualitative	Rheumatoid arthritis	86	 Age: 58.3 (13) to 59.6 (12.4) Female: 82.5% to 87% Ethnicity: 67.6–71.7% White, 13–18.9% Black, 15–26.1 Hispanic, 13.5–15.2 Other 	Observation tool for specific aspects of shared decision making
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Authors Year of publica- tion	Country	Study design	Condition(s)	Sam- ple size	Characteristics of participants Age=mean (SD) [Range]	SDM (instrument) measures
				(n)	Information on marital status,	
Hurley et al. 2020 [62]	United States of America	Quantitative	Hip or knee OA	5751	education level, employment, health insurance • Age: 58.2 to 60.6 • Female: 54.26% to 64.16% • Ethnicity: 65.9–79.1% White, 2.9–11.5% Hispanic or Latino, 7.1–9% Black or African American, 9.1–13.6% Non-	Record on having undergone arthroplasty
Hurley et al.	United States of	Quantitative	Hip or knee OA	1838	hispanic/non-white Information on marital status, health insurance, emotional distress Age: 58.5 (10.1) to 59.3 (9.6)	Item on treatment preferences
2020b [63]	America				 Female: 56.1% to 64.7% Ethnicity: 73.3–82.8% White, 17.1–26.7% Non-white/Other Information on marital status, education 	Item on decision making stage
Hutyra et al. 2019 [64]	United States of America	Quantitative	Anterior shoulder dislocation	199	 Age: 23.56 (5.27) [18–35] Female: 23% Ethnicity: NR Information on marital status, education, employment, income, health insurance 	 Decisional Conflict Scale Patient Activation Measure Analysis on treatment alignment with evidence-based treatment Item on stage of decision making Information on awareness of making a preference-sensitive decision Questionnaire on knowledge
Ibrahim et al. 2013 [66]	United States of America	Mixed methods	Knee OA	639	 Age: 60.70 (9.27) to 61.35 (8.73) Female: 6% to 7% Ethnicity: 100% Black Information on marital status, education level, employment, income, literacy 	retention Knowledge questions total knee replacement Items on importance Items on values and goals Items on summarizing pros and cons Questions on discussion of knee pain with primary care provider Items on readiness
Ibrahim et al. 2017 [65]	United States of America	Quantitative	Knee OA	336	 Age: 59.1 (7.2) Female: 69.9% Ethnicity: 100% Black Information on marital status, education level, employment, income 	Item on confidence Record on receiving total knee replacement Item on willingness to undergo surgery if recommended by the surgeon
Isaacs et al. 2013 [67]	United States of America	Quantitative	Orthopaedic injuries	111	Age: 73 (7) Female: 64% Ethnicity: 66% White, 34% African American Information on education level	Items on components of shared decision making Item on patient satisfaction with the treatment choice
Jayaku-mar et al. 2021 [68]	United States of America	Quantitative	Knee OA	129	 Age: 62.59 (8.85) to 62.62 (7.81) [45–89] Female: 62% to 67% Ethnicity: 32–41% White, 33–35% Hispanic or Latino, 16–18% Black or African American, 10–15% Asian Information on education level, employment, health insurance, emotional distress 	Knee-Decision Quality Instrument CollaboRATE Item on patient satisfaction with consultation Duration of consultations Analysis on total knee replacement rate Item on treatment concordance
Kane et al. 2023 [69]	United States of America	Mixed methods	Dupuytren contracture	30	 Age: 69 (8) Female: 17% Ethnicity: 97% White, 3% Black Information on education level, employment, income 	9-item Shared Decision Making Questionnaire Questions on satisfaction Item on support Questions on decision making process
Kearing et al. 2016 [70]	United States of America	Quantitative	Spinal stenosis	168	Age: 66.6 (9.7) to 66.7 (9.7)Female: 47% to 50%Ethnicity: 98% White	 Decisional Conflict Scale Knowledge questionnaire about treatment options (continued on next page)

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Authors Year of publica- tion	Country	Study design	Condition(s)	Sam- ple size (n)	Characteristics of participants Age=mean (SD) [Range]	SDM (instrument) measures
					Information on marital status, education level, employment, literacy	Item on stage of decision making
Kim et al. 2021 [71]	South Korea	Quantitative	Distal radius fractures	49	• Age: 55.7 (14.9) to 58.6 (8.4) • Female: 83.3% to 96% • Ethnicity: NR	Decisional Conflict Scale
Kjeken et al. 2006 [72]	Norway	Quantitative	Rheumatoid arthritis or ankylosing spondylitis	1193	 Age: 59.6 (15.6) Female: 74% Ethnicity: NR Information on education level, employment 	Items on received information Items on involvement in decisions Item on satisfaction with care Items on unmet health care needs
Kleiss et al. 2021 [73]	United States of America	Quantitative	Upper-extremity conditions (Trigger finger, Carpal tunnel syndrome, Thumb OA, Wrist ganglion, de Quervain tenosynovitis, Lateral epicondylitis, Distal radius fracture, Olecranon bursitis, Scaphoid fracture, Radial head fracture, Mallet fracture, Dupuytren disease)	147	 Age: 55 (14) [18–84] Female: 67% Ethnicity: 69% White, 31% non-White Information on marital status, education level, employment 	Decision Regret Scale Item on treatment choice Item on satisfaction with the visit
Knutsson et al. 2022 [74]	Sweden	Quantitative	Lumbar spine conditions (postsurgery)	209	Age: 64 (14)Female: 54%Ethnicity: NR	Items on shared decision making Item on overall satisfaction
Kravitz et al. 2018 [75]	United States of America	Quantitative	Musculoske-letal pain	215	 Age: 55.5 (11.1) Female: 47% Ethnicity: 74% White, 13% Black or African American, 11% Latino, 6% Asian, 8% Other Information on marital status, education level, employment 	with care Trust in Physician Scale Pain Medication in Primary Care Patient questionnaire Pain Treatment Satisfaction Scale Consumer Assessment of Healthcare Providers and System survey
Kunne-man et al. 2018 [76]	Netherlands	Qualitative	Osteoporosis	100	Age: 58 (13.2)Female: 50%Ethnicity: NRInformation on education level	OPTION Scale
Lai et al. 2021 [77]	United States of America	Quantitative	Displaced diaphyseal clavicle fractures	41	 Age: 39 (18) to 44 (15) Female: 13% to 16% Ethnicity: 60-75% White, 20-22% Asian, 20-22% Hispanic Information on education level, employment, health insurance 	Decisional Conflict Scale Record of treatment choice
LeBlanc et al. 2015 [78]	United States of America	Mixed methods	Osteopenia and osteoporosis	77	 Age: 66 (10) to 69 (8) Female: 100% Ethnicity: NR Information on education level, income, numeracy 	 Decisional Conflict Scale OPTION Scale Knowledge questionnaire about osteoporosis and treatment options Analysis on primary adherence Analysis on secondary adherence Duration of encounters Information on decision to start bisphosphonates
Li et al. 2014 [79]	Canada	Quantitative	Rheumatoid arthritis	30	 Age: 54.9 (14.9) Female: 76.7% Ethnicity: NR Information on marital status, education level, employment, income 	Decisional Conflict Scale Effective Consumer Scale Methotrexate in Rheumatoid Arthritis Knowledge test
Li et al. 2018 [80]	Canada	Quantitative	Rheumatoid arthritis	50	Age: 49.6 (12.2) Female: 80% Ethnicity: NR Information on marital status, education level, income	Decisional Conflict Scale (low literacy) Medication Education Impact Questionnaire Partners in Health Scale
Lofland et al. 2017 [81]	United States of America	Quantitative	Rheumatoid arthritis or psoriatic arthritis	204	 Age: 51 (11.3) to 51.3 (10.7) Female: 68.2% to 82.7% Ethnicity: 86.7–88.3% White, 6.7–7.8% Black or African American, 6.7–7.8% Hispanic, 1.6–2.7% Asian, 1.6–2.7% American Indian or Alaskan native, 2.3–4% Other 	9-item Shared Decision Making Questionnaire Patient Activation Measure Morisky Medication Adherence Scale
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Authors Year of publica- tion	Country	Study design	Condition(s)	Sam- ple size	Characteristics of participants Age=mean (SD) [Range]	SDM (instrument) measures
				(n)		
Lopez-Olivo et al. 2020 [82]	United States of America	Quantitative	Osteoporosis	225	 Age: 63.9 (8.5) Female: 100% Ethnicity: 45.3% White, 32.9% Hispanic, 15.6% Black or African American, 6.2% Other Information on marital status, education level, employment, literacy 	 Decisional Conflict Scale (low literacy) Osteoporosis Patient Knowledge Questionnaire Effective Consumer Scale
Mahlich et al. 2019 [83]	Japan	Quantitative	Rheumatoid Arthritis	500	 Age: 54.28 (10.02) Female: 67% Ethnicity: NR Information on marital status, education level, employment, income, emotional distress 	Items on preferences for shared decision making Item on satisfaction with treatment Treatment preference fit index
Mainz et al. 2022 [84]	Denmark	Mixed methods	Anterior cruciate ligament injury	50	Age: 27.6 [24.6–30.8]Female: 47%Ethnicity: NR	9-item Shared Decision Making Questionnaire Question on experience of shared decision making
Mangla et al. 2019 [85]	United States of America	Quantitative	Hip or knee OA	58	Age: 63 (9) to 64 (9)Female: 49% to 56%Ethnicity: NRInformation on literacy	Knee-Decision Quality Instrument Hip-Decision Quality Instrument
Marshall et al. 2023 [86]	Canada	Quantitative	Knee OA	140	 Age: 64.3 (8.7) to 64.4 (7.8) Female: 47.2% to 65.2% Ethnicity: NR Information on emotional distress 	Decision Regret Scale Item on patient expectations about knee replacement post-surgery Items on satisfaction with results of knee replacement
Martin et al. 2017 [87]	United States of America	Quantitative	Rheumatoid arthritis	399	 Age: 64.15 (12.79) to 64.92 (11.58) Female: 64.3% to 70% Ethnicity: 5.5–6.8% Minority Information on education level, income 	Decisional Conflict Scale Knowledge questionnaire about etanercept
Mathijs-sen et al. 2020 [88]	Netherlands	Qualitative	Rheumatoid arthritis	168	• Age: 61.2 (11.4) • Female: 69% • Ethnicity: NR • Information on education level	OPTION Scale
Meade et al. 2015 [89]	Australia	Quantitative	Rheumatoid arthritis	144	Age: 30.43 (5.07) to 31.26 (4.26) Female: 100% Ethnicity: NR Information on marital status, education level, emotional distress	Decisional Conflict Scale Pregnancy in Rheumatoid Arthritis Questionnaire
Montori et al. 2011 [90]	United States of America	Mixed methods	Osteoporosis	100	 Age: [50–84] Female: 100% Ethnicity: NR Information on education level, income 	 Decisional Conflict Scale OPTION Scale Trust in Physician Scale Knowledge questionnaire Information on satisfaction with knowledge transfer Item medication adherence Analysis on medication adherence Analysis on persistence
Nota et al. 2014 [92]	Netherlands	Quantitative	Rheumatoid arthritis, psoriatic arthritis, or ankylosing spondylitis	519	 Age: 56 (12) Female: 59% Ethnicity: NR Information on marital status, education level, employment, income 	Item on satisfaction with decision making process
Nota et al. 2016 [91]	Netherlands	Quantitative	Rheumatoid arthritis, psoriatic arthritis, or ankylosing spondylitis	281	 Age: 54 (15) to 55 (13) Female: 61% to 65% Ethnicity: NR Information on marital status, education level, employment 	 Control Preference Scale Satisfaction With Decision scale Beliefs about Medicines Questionnaire Morisky Medication Adherence Scale Satisfaction with Information about Medicines Scale Decision Evaluation Scales Cologne Patient Questionnaire

Authors Year of publica- tion	Country	Study design	Condition(s)	Sam- ple size (n)	Characteristics of participants Age=mean (SD) [Range]	SDM (instrument) measures
Oakley et al. 2006 [93]	United Kingdom	Quantitative	Osteoporosis	33	Age: 77 [61–90]Female: 100%Ethnicity: NR	Questionnaire on satisfaction with decision and decision making process Decisional Conflict Scale Beliefs about Medicines Questionnaires Satisfaction with Information about Medicines Scale Medication Adherence Report Scale
Pablos et al. 2020 [94]	Spain	Quantitative	Rheumatoid arthritis	54	Age: 58.82 (12.85)Female: 90.38%Ethnicity: NRInformation on marital status,	Analysis on compliance Decisional Conflict Scale Preparation for Decision Making scale Decision Self-Efficacy Scale
Patel et al. 2014 [95]	United Kingdom	Quantitative	Non-specific low back pain	148	education level • Age: 46.9 (13.8) to 48.8 (16.7) • Female: 65.1% to 67.1% • Ethnicity: 83.5–88.9% White, 4.8–8.2% Asian or Asian British, 3.2–3.5% Mixed, 3.2–3.5 Black or Black British, 0–1.2% Chinese • Information on employment,	Satisfaction With Decision scale Item on satisfaction with treatment Item on satisfaction with decision
Reilly et al. 2023 [96]	United States of America	Quantitative	Knee OA	20	emotional distress • Age: 71 (13.5) • Female: 5% • Ethnicity: 90% White, 5% Asian/Pacific Islander	Knee-Decision Quality Instrument Items on overall experience with decision making process
Rivero- Santana et al. 2021 [97]	Spain	Quantitative	Knee OA	193	 Information on education level Age: 66.79 (8.42) Female: 72.02% Ethnicity: NR Information on education level 	Decisional Conflict Scale Knee-Decision Quality Instrument Decision Regret Scale Items on satisfaction with the decision making process Information on having undergone surgery
Sanders et al. 2022 [98]	Netherlands	Mixed methods	Non-chronic low back pain	176	Age: 46.77 (13.16)Female: 53.8%Ethnicity: NR	Item on treatment preference OPTION Scale Item on patient-reported shared decision making
Scoville et al. 2011 [99]	United States of America	Qualitative	Osteoporosis	18	 Information on education level Age: 70.6 (9.4) Female: 100% Ethnicity: NR 	 Observation grid on the reasons women present when expressing hesitation about initiation of bisphosphonates and how clinicians react
Sepucha et al. 2011 [105]	United States of America	Validation	Hip or knee OA	509	 Age: 62.7 (9.6) to 66.1 (9.49) Female: 56% to 59.1% Ethnicity: 95.5% White Information on marital status, education level, income 	Knee-Decision Quality Instrument Hip-Decision Quality Instrument
Sepucha et al. 2012 [104]	United States of America	Validation	Lumbar herniated disc	341	 Age: 44 (8.6) to 48 (9.6) Female: 45% to 54% Ethnicity: 72.5–94% White, 2–20% Black, 2–13% Hispanic, 0.6–3% Asian, 1–8.5% Other Information on education level, income 	Herniated Disc-Decision Quality Instrument
Sepucha et al. 2013 [102]	United States of America	Quantitative	Hip or knee OA	382	 Age: 62.7 (9.6) Female: 55.8% Ethnicity: 93.9% White, 97.6% Hispanic Information on education level 	Knee-Decision Quality Instrument Hip-Decision Quality Instrument Item on decision regret Items on decision process Item on decision confidence
Sepucha et al. 2017 [100]	United States of America	Quantitative	Hip or knee OA, lumbar spinal stenosis, or lumbar herniated disc	649	 Age: 62.7 (13.1) to 63.8 (11.9) Female: 50% to 52.9% Ethnicity: 91–91.4% Nonhispanic white, 2.4–3.1% Black, 3.1% Other or multiple, 0.9–1.5% Hispanic Information on education level 	Knee-Decision Quality Instrument Hip-Decision Quality Instrument Herniated Disc-Decision Quality Instrument (continued on next page)

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Authors Year of publication	Country	Study design	Condition(s)	Sam- ple size (n)	Characteristics of participants Age=mean (SD) [Range]	SDM (instrument) measures
Sepucha et al. 2018 [103]	United States of America	Quantitative	Hip or knee OA, lumbar spinal stenosis, or lumbar herniated disc	543	 Age: 63.9 (12.1) Female: 52.7% Ethnicity: 92.4% Non-hispanic white, 2.9% Other or multiple, 2.4% Black, 2.6% Hispanic Information on education level 	Spinal Stenosis-Decision Quality Instrument CollaboRATE Shared Decision-Making Process Record on having undergone surgery Knee-Decision Quality Instrument Hip-Decision Quality Instrument Herniated Disc-Decision Quality Instrument Spinal Stenosis-Decision Quality Instrument Decision Regret Scale Items on patient's satisfaction with how their treatment
Sepucha	United States of	Quantitative	Hip or knee OA	1124	• Age: 65 (10)	 Items on patient's satisfaction with their current pain and symptoms Record on treatment received Knee-Decision Quality
et al. 2019 [101]	America				Female: 57% Ethnicity: 89% Non-hispanic white Information on education level, literacy, health incurance.	Instrument • Hip-Decision Quality Instrument • Shared Decision-Making Process
Sepucha et al. 2022 [106]	United States of America	Quantitative	Hip or knee OA	845	literacy, health insurance • Age: 65 (9) • Female: 58% • Ethnicity: 93% Non-hispanic white • Information on education level, health insurance	Knee-Decision Quality Instrument Hip-Decision Quality Instrument Decision Regret Scale Item on satisfaction with treatment Item on satisfaction with their current pain Analysis on informed, patient-
Shaw et al. 2021 [107]	Switzerland	Quantitative	Rheumatoid arthritis, psoriatic arthritis, or axial spondyl-arthritis	2111	 Age: 46.6 (12) to 50.9 (13.1) Female: 55% to 57% Ethnicity: NR Information on education level, 	centred decision Record on treatment received CollaboRATE Information on satisfaction with shared decision making
Shirley et al. 2015 [108]	United States of America	Quantitative	Neuro-muscular scoliosis (parents reported the information)	11	employment • Age: 12.2 [8–17] • Female: NR • Ethnicity: NR	SURE test Knowledge questionnaire about neuromuscular scoliosis treatment Items on parent's satisfaction
Shue et al. 2016 [109]	United States of America	Quantitative	Hip or knee OA	147	 Age: 61 (11) Female: 53% Ethnicity: 50% White, 33% African American, 12% Hispanic, 4% Asian, 1% Other Information on education level, health insurance 	with shared decision making Knowledge questionnaire about hip or knee OA disease progression and total hip or knee arthroplasty Item on satisfaction regarding education and knowledge Items on decision making participation Item on treatment preference Item on stage of decision
Simon et al. 2012 [110]	Germany	Quantitative	Acute low back pain	2480	 Age: 45.34 (12.99) to 45.81 (12.71) Female: 52% to 52.4% Ethnicity: NR Information on marital status, education level, native language, emotional distress 	making Decisional Conflict Scale Decision Regret Scale Preparation for Decision Making scale Perceived Involvement in Care Scale Knowledge questionnaire Item on treatment adherence Item on patient preference for participation

Authors Year	Country	Study design	Condition(s)	Sam-	Characteristics of participants	SDM (instrument) measures
of publica- tion				ple size (n)	Age=mean (SD) [Range]	
Small-wood et al. 2017 [111]	United States of America	Quantitative	Osteopenia or osteoporosis	50	Age: 67.8 to 68.8 Female: 100% Ethnicity: 98% White, 2% African American, 4% Hispanic Information on marital status, education level, employment, income, health insurance	Decisional Conflict Scale Preparation for Decision Making scale Information on treatment decision Information on patient- reported decision making
Stacey et al. 2014 [112]	Canada	Quantitative	Knee OA	137	 Age: 67.1 (10.85) to 67.3(12.16) Female: 64.8% to 72.5% Ethnicity: NR Information on education level, employment 	Knee-Decision Quality Instrument Preparation for Decision Making scale SURE test Decision quality analysis
Stacey et al. 2016 [113]	Canada	Quantitative	Hip or knee OA	334	 Age: 66.1 (9.8) to 66.9 (9.8) Female: 53.4% to 61.7% Ethnicity: NR Information on education level, employment, income, native language 	Knee-Decision Quality Instrument Hip-Decision Quality Instrument Preparation for Decision Making scale SURE test Decision quality analysis Analysis on realistic expectation Surgical rate
Sumpton et al. 2022 [114]	Australia	Qualitative	Psoriatic arthritis	25	 Age: [27–79] Female: 44% Ethnicity: 72% Australia/New Zealand, 16% Asia/Pacific, 8% Americas, 4% Europe Information on education level, employment 	Questions on values Questions on satisfaction with received information Questions on relationship and communication with clinicians Questions on confidence with current understanding Questions on control when making a decision Questions on decision making process
Torrente- Jimenez et al. 2022 [115]	Spain	Quantitative	Knee OA	193	 Age: 66.8 (8.42) Female: 72% Ethnicity: NR Information on education level 	Decisional Conflict Scale Knowledge questionnaire about OA and total knee replacement Information on the importance given to certain characteristics and potential outcomes of OA treatments Items on satisfaction with decision making process Item on treatment preference Item on having undergone surgery
Tutuha- tunewa et al. 2017 [116]	Netherlands	Quantitative	Midshaft clavicle fracture	278	Age: 39.7 to 42.4 [23.6–55.8]Female: 14.1% to 22%Ethnicity: NR	Items on overall satisfaction with care
Valentine et al. 2021 [117]	United States of America	Quantitative	Hip or knee OA, lumbar spinal stenosis, or lumbar herniated disc	168	 Age: 65 (11) Female: 52% Ethnicity: 93% White non-Hispanic 	Knee-Decision Quality Instrument Hip-Decision Quality Instrument Herniated Disc-Decision Quality Instrument Spinal Stenosis-Decision Quality Instrument Shared Decision-Making Process Analysis on informed, patient-centred decision
van Dijk et al. 2021 [118]	Netherlands	Quantitative	Hip or knee OA	131	 Age: 66 (10) to 68 (11) Female: 50% to 54% Ethnicity: NR Information on marital status, education level, employment 	 Decisional Conflict Scale Items on satisfaction with the given information, the clinic, and the physician Knowledge questionnaire about treatment options and risks Information on stage of decision making (continued on next page)

Authors Year of publica- tion	Country	Study design	Condition(s)	Sam- ple size (n)	Characteristics of participants Age=mean (SD) [Range]	SDM (instrument) measures
Volk-mann et al. 2015 [119]	United States of America	Quantitative	Knee OA	111	 Age: 70 (9.6) to 72 (8.2) Female: 63% Ethnicity: NR Information on marital status, 	Information on treatment preference Information whether patient had made their definitive decision after the first visit Decisional Conflict Scale Item on decision readiness
Weng et al. 2007 [120]	United States of America	Quantitative	Knee OA	64	education level • Age: NR • Female: NR • Ethnicity: 51.5% African American, 48.4% Caucasian	Decisional Conflict Scale Item on decision readiness Item on stage of decision making Item on willingness to consider total knee replacement Item on beliefs about the effectiveness of joint replacement
Wilkens et al. 2019 [121]	United States of America	Quantitative	Trapezio-metacarpal arthritis	90	 Age: 65 (1.3) to 65 (1.5) Female: 49% to 51% Ethnicity: 48–52% White Information on marital status, education level, employment 	Decisional Conflict Scale Decision Regret Scale Consultation and Relational Empathy Scale Information on treatment choice Item on satisfaction with the visit Item on overall treatment satisfaction Analysis for change of treatment
Youm et al. 2015 [122]	United States of America	Quantitative	Hip or knee OA	123	 Age: 62.4 (11.4) to 63.8 (9.31) [19–85] Female: 54% Ethnicity: 74% Non-hispanic, 7% Hispanic Information on education level, income, health insurance 	Knee-Decision Quality Instrument Hip-Decision Quality Instrument Analysis on informed, patient centred decision Stage of Decision Making scale Information on treatment choice
Zadro et al. 2022 [123]	Australia, New Zealand, United Stated of America, United Kingdom, Canada	Quantitative	Subacromial pain syndrome	409	 Age: 41.3 (10) Female: 44.2% Ethnicity: NR Information on education level, employment, health insurance 	Decisional Conflict Scale Item on treatment intention Knowledge questionnaire about options Analysis for informed choice Items on attitudes towards surgery

SD: Standard deviation SDM: Share decision making

OA: osteoarthritis NR: Not reported

FAPI: Fragebogen zur Arzt-Patienten-Interaktion

OPTION Scale: Observing Patient Involvement in Decision Making instrument

MASRI: Medication Adherence Self-Report Inventory

Appendix B. Characteristics of the measurement instruments with any measurement properties reported by the included studies

Measure-ment instrument	Extracted construct	Condition(s) in which it was used	Type of measu- re	Number of items and subscales	Score, cut-off and interpretation	Information on measurement properties
Decisional Conflict Scale	Decisional conflict [120, 123] Aspect of the decision making process [30] Experiences quality of the decision [34,87,90, 111] Perceptions of being uncertain, uninformed, unsupported, or unclear	 Knee OA (n=9) [34,43, 44,46,97,115,118-120] Hip OA (n=3) [34,46, 118] Trapeziometacarpal arthritis (n=1) [121] Lumbar herniated disc (n=1) [26] Lumbar degenerative diseases (n=1) [41] 	PROM	16 items 3 subscales (healthcare consumers' uncertainty in making a health-related decision, factors contributing to the uncertainty, healthcare consumers' perceived effective decision	0−100 [26,43,123] Higher score indicating greater decisional conflict [26,123] Total score ≤ 25: tend to make decisions [42,43,87,94] Total score > 37.5: tend to delay decisions or to feel unsure about	Validity: • Validated: - no information in which populations [41,46,118] • Discriminant validity: - no information in which populations [42,87,119,120,123] Reliability:
						(continued on next page)

constructs:
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Measure-ment instrument	Extracted construct	Condition(s) in which it was used	Type of measu- re	Number of items and subscales	Score, cut-off and interpretation	Information on measurement properties
	as to values to be considered [44,46,94] • Patient's uncertainty in making a given health-related decision [41,55,71,121] • Personal perceptions of: (1) uncertainty in health-related decision making; (2) factors contributing to the uncertainty and; (3) the perceived effectiveness of decision making [89].	 Spinal stenosis (n=1) [70] Rheumatoid arthritis (n=7) [50,51,53,79,87,89,94] Osteopenia or osteoporosis (n=5) [42,78,90,93,111] Fibromyalgia (n=2) [30,58] Carpal tunnel syndrome (n=1) [55] Anterior shoulder dislocation (n=1) [64] Distal Radius Fractures (n=1) [71] Displaced diaphyseal clavicle fractures (n=1) [77] Acute low back pain (n=1) [110] Subacromial pain syndrome (n=1) [123] 		making subscales) [34, 42,79,118,121] 4 subscales (Informed, Values Clarity, Support, and Effective Decision subscales) [87] 5 subscales (being informed, values clarity, support, uncertainty, and effective decision-making subscales) [41, 43,53,64,89,94,97]	implementation [43,87, 94] • Total score > 38: tend to delay decisions [42] • Scores ≤ 2.0: no difficulty in decision making and implementation [93] • Scores ≥ 2.5: decision delay [93]	Reliable: no information in which populations [41,42,46,118] Internal consistency: in rheumatology [53] in women with rheumatoid arthritis [89] in low back pain [110] no information in which populations [123] Test-retest reliability: no information in which populations [89,121, 123] Longitudinal construct validity: Sensitive to change: no information in which populations
Decisional Conflict Scale (Low literacy)	Decisional conflict [25, 82] Quality of decision [29] Perceived uncertainty in choosing options, factors contributing to uncertainty, and effective decision making [59,80]	 Knee OA (n=2) [25,59] Hip OA (n=1) [25] (Allen 2016) Rheumatoid arthritis (n=2) [29,80] Osteoporosis (n=1) [82] 	PROM	• 10 items [25,59,80] • 4 subscales [25,59,80] 0-5 subscales [82]	 0-100 [25,59,80,82] Higher scores indicating greater decisional conflict [25,59,80] 0: being extremely well-informed and clear, and 100: being extremely uninformed or unclear [82]. Total score < 25: tend to make decisions [59,80] Total score > 37.5: tend to delay decisions [59,80] 	[42] Validity: Discriminant validity: no information in which populations [25,59] Correlation with other constructs: no information in which populations [59] Reliability: Internal consistency: no information in which populations [25,59,80,82] Test-retest reliability:
SURE test	Decisional conflict [27, 33,108] Patient's perception of feeling sure, informed, supported, and clear about what mattered most [39,112,113].	 Knee OA (n=5) [27,33, 39,112,113] Hip OA (n=2) [39,113] Lumbar herniated disc (n=1) [39] Lumbar spinal stenosis (n=1) [39] Juvenile Idiopathic Arthritis (n=1) [38] Neuromuscular scoliosis (parents made the decision) (n=1) [108] 	PROM	• 4 items [27,33,38,39, 108,112]	Response of "yes" to all 4 items indicates no uncertainty [38] Patients who answered "no" to any SURE test item were experiencing decisional conflict [108] Cut-off of 3 or less identifies clinically significant decisional conflict [39]	- no information in which populations [80,82] Validity: • Validated: - no information in which populations [108] • Discriminant validity: - in people with hip or knee OA [39] • Predictive validity: - in people with hip or knee OA [39] - in people with lumbar herniated disc or lumbar spinal stenosis [39] • Construct validity: - in people with hip or knee OA [39] - in people with hip or knee OA [39] - in people with lumbar herniated disc or lumbar spinal stenosis [39] - Correlation with other constructs:

changes in knowledge (continued on next page)

Extracted construct	Condition(s) in which it was used	Type of measu- re	Number of items and subscales	Score, cut-off and interpretation	Information on measurement properties
					 no information in which populations [112,113] <i>Reliability:</i> Internal consistency:
How well the intervention helped them with various aspects of decision making [25,94] Patients' perception of the usefulness of the intervention in preparing them to communicate with their physician [52,59] Preparation for decision making [59,110] Patients' perceptions of the decision making process [112]	 Knee OA (n=4) [25,59, 112,113] Hip OA (n=2) [25,113] Rheumatoid arthritis (n=1) [94] Osteopenia or osteoporosis (n=1) [111] Pain involving one or both knees (n=1) [52] Acute low back pain (n=1) [110] 	PROM	• 10 items [25,59, 111-113] • 11 items [52,59]	 0–100 [25,59,110,111] Higher indicating greater preparation [25, 59,110,111] 	 no information in which populations [39,112, 113] Validity: Discriminant validity: no information in which populations [52,59,112,113] Correlation with other constructs: no information in which populations [59] Reliability: Internal consistency: no information in which populations [52,59,110,112]
Knowledge about health conditions and treatment options (exclusively knowledge subscale) [25,27,40,96,97] Patient's knowledge and readiness to decide [48] Decision process (exclusively for talking with health care providers subscale) [68] The extent to which patients were informed and received their preferred treatment [100]. Patient's decision quality [113] Determine whether or not a decision was informed and patient-centred [117,122] Understanding of key facts about the treatment options [59]	• Knee OA (n=20) [25, 27,33,40,48,59,68,85, 96,97,100-103,105, 106,112,113,117,122]	PROM	 4 items (knowledge subscale) [112] 5 items [40,59] (knowledge subscale) 6 items (short version) (knowledge, treatment preference subscales) [101,106,117] 7 items (knowledge subscale) [97] 9 items (knowledge subscale) [25] 9 items (knowledge, goals and concerns, treatment preference subscales) [85,100,103] 13 items (knowledge, readiness to decide, influences on patient's decisions subscales) [48] 16 items (knowledge, goals, and concerns subscales) [96,105] 25 items (knowledge and values subscales) [113] 	 0–100 [25,40,59,85, 100,101] Higher scores indicating greater knowledge [25, 40,59,85,100,101] Knowledge thresholds were based on recommendations from the DQI scoring guides [117]. 	Validity: • Validated: - no information in which populations [27,33,40,100,101, 103,106,117] • Discriminant validity: - no information in which populations [102,113] - in people with knee OA [105] • Predictive validity: - no information in which populations [25,102,113] - in people with knee OA [105] • Content validity: -no information in which populations [25,102,113] - in people with knee OA [105] Reliability: • Reliable: - no information in which populations [25,100,113] • Test-retest reliability: - no information in which populations [101-103,106,117] - in people with knee OA [105] Responsiveness: • Sensitivity to change: - no information in which populations [101,103,106,117]
	How well the intervention helped them with various aspects of decision making [25,94] Patients' perception of the usefulness of the intervention in preparing them to communicate with their physician [52,59] Preparation for decision making [59,110] Patients' perceptions of the decision making process [112] Knowledge about health conditions and treatment options (exclusively knowledge subscale) [25,27,40,96,97] Patient's knowledge and readiness to decide [48] Decision process (exclusively for talking with health care providers subscale) [68] The extent to which patients were informed and received their preferred treatment [100]. Patient's decision quality [113] Determine whether or not a decision was informed and patient-centred [117,122] Understanding of key facts about the	• How well the intervention helped them with various aspects of decision making [25,94] • Patients' perception of the usefulness of the intervention in preparing them to communicate with their physician [52,59] • Prepration for decision making [59,110] • Patients' perceptions of the decision making process [112] • Knowledge about health conditions and treatment options (exclusively knowledge subscale) [25,27,40,96,97] • Patient's knowledge and readiness to decide [48] • Decision process (exclusively for talking with health care providers subscale) [68] • The extent to which patients were informed and received their preferred treatment [100]. • Patient's decision quality [113] • Determine whether or not a decision was informed and patient-centred [117,122] • Understanding of key facts about the	• How well the intervention helped them with various aspects of decision making [25,94] • Patients' perception of the usefulness of the intervention in preparing them to communicate with their physician [52,59] • Preparation for decision making [59,110] • Patients' perceptions of the decision making [59,110] • Patients' perceptions of the decision making process [112] • Knowledge about health conditions and treatment options (exclusively knowledge subscale) [25,27,40,96, 97] • Patient's knowledge and readiness to decide [48] • Decision process (exclusively for talking with health care providers subscale) [68] • The extent to which patients were informed and received their preferred treatment [100]. • Patient's decision quality [113] • Determine whether or not a decision was informed and patient-centred [117,122] • Understanding of key facts about the	• How well the intervention helped them with various aspects of decision making [25,94] • Patients' perception of the decision making [59,110] • Preparation for decision making [59,110] • Patients' perceptions of the decision making [59,110] • Patients' perceptions of the decision making sprocess [112] • Knowledge about health conditions and treatment options (exclusively knowledge ausbascale) [25,27,40,96,97] • Patient's knowledge and readiness to decide [48] • Decision process (exclusively for talking with health care providers subscale) [68] • The extent to which patients were informed and received their preferred treatment [100]. • Patient's decision quality [113] • Determine whether or not a decision was informed and a patient-centred [117,122] • Understanding of key facts about the treatment options [59]	* How well the intervention helped them with various aspects of decision making [25,94] * Patients' perception of the usefulness of the intervention in perparting them to communicate with their physician [32,59] * Preparation for decision making [25,94] * Acute low back pain (n=1) [110] * Patients' perceptions of the decision making [25,94] * Knowledge about health conditions and treatment options (exclusively knowledge subscale) [25,27,40,96,97] * Patient's knowledge and readiness to decide [48] * Decision process (exclusively for talking with health care providers subscale) [68] * The exent to which patients were informed and received their perferred treatment [100]. * Patient's decision quality [113] * Determine whether or not a decision was informed and patient centred [117,122] * Understanding of key facts about the treatment options [59]

Measure-ment instrument	Extracted construct	Condition(s) in which it was used	Type of measu- re	Number of items and subscales	Score, cut-off and interpretation	Information on measurement properties
						and concordance scores are 10% [85] Feasibility:
Hip-Decision Quality Instrument	Knowledge about health conditions and treatment options (exclusively knowledge subscale) [25,40] The extent to which patients were informed and received their preferred treatment [100]. Patient's decision quality [113] Determine whether or not a decision was informed and patient-centred [117,122]	• Hip OA (n=12) [25,40, 85,100-103,105,106, 113,117,122]	PROM	 5 items (knowledge subscale) [40] 6 items (short version) (knowledge, treatment preference subscales) [101,106,117] 9 items (knowledge subscale) [25] 9 items (knowledge, goals and concerns, treatment preference subscales) [85,100,103] 16 items (knowledge, goals, and concerns subscales) [105] 25 items (knowledge and values subscales) [113] 	 0-100 [25,40,85,100, 101] Higher scores indicating greater knowledge [25, 40,85,100,101] Knowledge thresholds were based on recommendations from the DQI scoring guides [117]. 	Evidence of Acceptability [100-103, 105,106,117] Evidence of feasibility [101-103,105,106,117] Evidence of feasibility [101-103,105,106,117] Validated:
						 no information in which populations [25,100,113] Test-retest reliability: no information in which populations [101-103,106,117] in people with hip OA [105] Responsiveness:
						• Sensitivity to change: - no information in which populations [101,103,106,117] Interpretability:
						• The minimal important changes in knowledge and concordance scores are 10% [85] Feasibility:
Herniated Disc- Decision Quality Instrument	Knowledge about health conditions and treatment options (exclusively knowledge subscale) [40] The extent to which patients were informed and received their	• Lumbar herniated disc (n=6) [26,40,100,103, 104,117]	PROM	 5 items (knowledge subscale) [40] 6 items (knowledge, treatment preference subscales) [117] 9 items (knowledge, goals and concerns, treatment preference subscales) [100,103] 	 0-100 [26,40,100,104] Higher scores indicating greater knowledge [40]. Each dimension has a separate total score ranging from 0-100, [0 = no knowledge or no involvement in the decision; 100 = best 	Evidence of Acceptability [100-103, 105,106,117] Evidence of feasibility [101-103,105,106,117] Validity: Validated:

(continued on next page)

Measure-ment instrument	Extracted construct	Condition(s) in which it was used	Type of measu- re	Number of items and subscales	Score, cut-off and interpretation	Information on measurement properties
	preferred treatment [26, 100]. • Determine whether or not a decision was informed and patient-centred [117]		·	• 19 items (knowledge and concordance subscales) [104]	possible knowledge or best possible involvement in the decision] [26].	 in people with herniated disc [104] Predictive validity: in people with herniated disc [104] Convergent validity: in people with herniated disc [104] Reliability:
						Reliable: no information in which populations [100] Test-retest reliability: no information in which populations [101,103,117] in people with herniated disc [105] Responsiveness:
						Sensitivity to change: no information in which populations [101,103,117] Feasibility:
Spinal stenosis-	 Knowledge about health 	Lumbar spinal stenosis	PROM	• 5 items (knowledge	• 0-100 [40,100]	 Evidence of Acceptability [100,101, 103,104,117] Evidence of feasibility [101,103,104,117] Validity:
Decision Quality Instrument	conditions and treatment options (exclusively knowledge subscale) [40] • The extent to which patients were informed and received their	(n=4) [40,100,103, 117]		subscale) [40] • 6 items (knowledge, treatment preference subscales) [117] • 9 items (knowledge, goals and concerns, treatment preference	Higher scores indicating greater knowledge [40].	• Validated: - no information in which populations [40,100,103,117] Reliability:
	preferred treatment [100]. Determine whether or not a decision was informed and patient-centred [117]			subscales) [100,103]		Reliable: no information in which populations [100] Test-retest reliability: no information in which populations [103,117] in people with herniated disc [105] Responsiveness:
						Sensitivity to change: no information in which populations [103,117] Feasibility:
ReproKnow	Reproductive	• Rheumatic diseases	PROM	• 10 items	• 0–10 [31]	 Evidence of Acceptability [100,103, 117] Evidence of feasibility [103,117] Validity:
	knowledge across a range of topical domains [31]	(n=1) [31]			10 indicating a perfect score on the assessment [31]	Content validity: -in women with rheumatic diseases [31] Known group validity: - in women with rheumatic diseases [31] Structural validity: (continued on next page)

Measure-ment instrument	Extracted construct	Condition(s) in which it was used	Type of measu- re	Number of items and subscales	Score, cut-off and interpretation	Information on measurement properties
						- in women with rheumatic diseases [31] Reliability:
						• Internal consistency: - in women with rheumatic diseases [31] Feasibility:
Methotre-xate	Knowledge about	Rheumatoid arthritis (a. 1) [70]	PROM	• 60 items [79]	• 0–60 [79]	• Evidence of feasibility [31] Reliability:
in rheumatoid arthritis knowledge test	methotrexate [79]	(n=1) [79]				• Internal consistency: - in people with rheumatoid arthritis [79]
						 Test-retest reliability: in people with rheumatoid arthritis [79]
Osteoporosis patient knowledge question- naire	Osteoporosis knowledge [82]	• Osteoporosis (n=1) [82]	PROM	• 17 items (20 items in the original version) [82]	 0-17 [82] Higher scores indicating superior knowledge [82] 	Validated: - assumed in people with osteoporosis [82] Reliability:
						• Reliable: - assumed in people with osteoporosis [82]
Pregnancy in rheumatoid arthritis question- naire	 Rheumatoid arthritis, pregnancy, and parenting knowledge [89] 	• Rheumatoid arthritis (n=1) [89]	PROM	• 39 items	0-39 [89]Higher scores indicating greater knowledge [89]	Reliability: • Internal consistency: - in women with rheumatoid arthritis [89]
Collabo-RATE	Patient involvement in the decision making process [26] Patient's perception of how much effort was made to help them understand their health issue, how much the provider listened to them about their health issue, and how much effort was made to include what matters most to the patient in choosing what to do next [39,100] Level of shared decision making [68] Shared decision making [107]	Knee OA (n=4) [27,39, 68,100] Hip OA (n=2) [39,100] Lumbar herniated disc (n=3) [26,39,100] Lumbar spinal stenosis (n=2) [39,100] Rheumatoid arthritis (n=1) [107] Juvenile idiopathic arthritis (n=1) [38] Lupus nephritis (n=1) [54] Psoriatic arthritis (n=1) [107] Axial spondylarthritis (n=1) [107]	PROM	• 3 items [26,27,38,39, 54,100,107]	 0-9 [26,38,39,54,100] Higher scores indicating more clinician effort to engage and involve the parent [26,38,39,54, 100] 	Validated:

Measure-ment instrument	Extracted construct	Condition(s) in which it was used	Type of measu- re	Number of items and subscales	Score, cut-off and interpretation	Information on measurement properties
						Sensitivity to change: no information in which populations [38]
Control Preference Scale	Perceived and/or preferred role in medical decision making [91] The extent of decision making control a patient preferred in treatment decisions [41]	 Knee OA (n=1) [27] Lumbar degenerative diseases (n=1) [41] Rheumatoid arthritis (n=1) [91] Psoriatic arthritis (n=1) [91] Ankylosing spondylitis (n=1) [91] 	PROM		 A score of 1 indicates a preference for full patient autonomy in decision making, a score of 5 corresponds to a preference for physicians making decisions [41]. 	Validity: Convergent validity: no information in which populations [41] Reliability: Internal consistency: no information in which populations
Trust in Physician Scale	• Trust [29,90] • Trust in physician [28]	 Rheumatoid arthritis (n=2) [28,29] Osteoporosis (n=1) [90] Musculoskeletal pain (n=1) [75] 	PROM	• 11 items	0–100, a score below the median (90.9) was considered to be suboptimal [28]	 [41] Validity: Validated: in people with rheumatoid arthritis
Interperso-nal Processes of Care	Physician-patient interactions (communication, patient-centred decision making, and physician interpersonal style) [45] Patient perception of communication around shared decision making [28]	• Rheumatoid arthritis (n=2) [28,29] • Systemic lupus erythematosus (n=1) [45]	PROM	 29 items [45] 3 subscales (communication, decision making, and interpersonal style) [45] 	Not clear	Validity: • Validated: - no information in which populations [28] - in socioeconomically and ethnically diverse populations of adults from general medicine practices [45] Reliability:
						• Reliable: - no information in which populations [28,45]
Medication adherence	Medication adherence [29]	• Rheumatoid arthritis (n=1) [29]	PROM	• 1 item [29]	A response of 1 or greater was considered non-adherent [29]	Validity: • Validated: - no information in which populations [29]
Satisfaction with decision scale	 The results of the decision making process [30,58] Patient satisfaction with health care decisions [41,53] Consistency with personal values subscale measures whether the 	 Lumbar degenerative diseases (n=1) [41] Rheumatoid arthritis (n=2) [53,91] Juvenile idiopathic arthritis (n=1) [53] Psoriatic arthritis (n=2) [53,91] Ankylosing spondylitis 	PROM	 2 items (consistency with personal values subscale) [91] 6 items [41,53,58] 	 Higher total scores denote higher satisfaction with a decision [41,53] Higher score indicating higher consistency with personal values [91] 	Validity: Construct validity: no information in which populations [41] Reliability: Reliable:
	decision meets personal preference measures whether the decision meets personal preferences [91] • Satisfaction with treatment [95]	(n=2) [53,91] Granulomatosis with polyangiitis (n=1) [53] Other rheumatic diseases (n=1) [53] Fibromyalgia [30,58] (n=2) Non-specific low back pain (n=1) [95]				 in patients with rheumatic diseases and their companions [53] Internal consistency: no information in which populations [41,91] in people with rheumatic diseases and their companions [53]
Question-naire on Doctor- Patient Interaction (FAPI)	Quality of physician–patient interaction from the patients' perspective (adequate imparting of information, involvement in medical decisions, and a feeling	• Fibromyalgia (n=1) [30]	PROM	• 14 items [30]	 1–5 [30] Higher score indicating higher quality [30] 	Validity: • Discriminant validity: - in patients from outpatient clinics for general internal medicine, diabetes, (continued on next page)

Measure-ment instrument	Extracted construct	Condition(s) in which it was used	Type of measu- re	Number of items and subscales	Score, cut-off and interpretation	Information on measurement properties
	of being taken seriously by the physician) [30]					rheumatology, and pain [30] Reliability:
						Internal consistency: in patients from outpatient clinics for general internal medicine, diabetes, rheumatology, and pain [30] Feasibility:
9-item Shared Decision Making Question- naire	 Perceived quality of the decision process [34,41] Patient's perceived involvement in shared decision making [47,81] 	 Knee OA (n=1) [34] Hip OA (n=1) [34] Lumbar degenerative diseases (n=1) [41] Rheumatoid arthritis 	PROM	• 9 items [34,41,47,84]	 0-100 [34,41,69,81] High score means high perceived level of shared decision making [34,41, 69,81,84] 	 Evidence of feasibility [30] Validity: Validated: no information in which populations
nenc	Perceived shared decision making [69,84]	 (n=1) [81] Juvenile idiopathic arthritis (n=1) [47] Psoriatic arthritis (n=1) [81] Dupuytren contracture (n=1) [69] 			69,81,84]	[41,47] • Construct validity: - no information in which populations [69] Reliability:
		• Anterior cruciate ligament injury (n=1) [84]				Reliable: no information in which populations [41] Internal consistency: no information in which populations [69]
Princess Margaret Hospital Satisfaction with Doctor Question-	• Satisfaction on patients' physician interaction [35]	• Nontraumatic painful conditions of the upper extremity (n=1) [35]	PROM	• 29 items [35]	Higher score reflects higher satisfaction with the doctor-patient inter- action [35]	Validity: • Validated: - in oncologic patients [35]
naire Informed Shared Decision- Making Scale	Competencies that physicians should pursue for informed shared decision making [35]	• Nontraumatic painful conditions of the upper extremity (n=1) [35]	PROM	• 16 items [35]	0-32 [35] A higher coding score indicates a greater level of informed shared decision making [35]	Validity: • Validated: - no information in which populations [35] Reliability:
						• Reliable: - no information in which populations [35]
Decision Regret Scale	 Distress or remorse after a decision [39,40,73, 103,106] Decisional regret [86,97, 110,121] 	 Knee OA (n=6) [39,40, 86,97,103,106] Hip OA (n=4) [39,40, 103,106] Trapeziometacarpal arthritis (n=1) [121] Lumbar herniated disc (n=3) [39,40,103] Lumbar spinal stenosis (n=3) [39,40,103] Upper-extremity conditions (n=1) [73] Acute low back pain (n=1) [110] 	PROM	• 5 items [39,40,73,86, 97,103,106,121]	 0-20 [106] 0-100 [39,40,73,97, 103,110,121] Higher scores indicate more regret [39,40,73, 97,103,106,110,121] 	Validity: • Validated: - no information in which populations [40,86,106] • Correlation with other constructs: - no information in which populations [103,121] • Measurement invariance: - no information in which populations [121]
						Reliability: • Reliable: (continued on next page)

Measure-ment instrument	Extracted construct	Condition(s) in which it was used	Type of measu- re	Number of items and subscales	Score, cut-off and interpretation	Information on measurement properties
Shared	• The extent of the	• Knee OA (n=5) [39,40,	PROM	• 4 items [39,40,101,	• 0-4 [39,40,101,117]	- no information in which populations [106] • Internal consistency: - no information in which populations [39,103] - in people with low back pain [110] Validity:
Decision- Making Process	interaction between the provider and patient that meet the standards of shared decision making [39] Patient involvement in decision making [40] The amount of shared decision making in the visit (including discussion of surgical procedures and nonsurgical options, the advantages and disadvantages of each, and patients' preferences) [100,101]	100,101,117] • Hip OA (n=5) [39,40, 100,101,117] • Lumbar herniated disc (n=4) [39,40,100,117] • Lumbar spinal stenosis (n=4) [39,40,100,117]		117] • 7 items [100]	 0-100 [100] Higher score indicating more involvement in the decision [39,40] Higher scores indicating more shared decision making [100,101,117] 	Discriminant validity: in people with hip or knee OA [39] Predictive validity: in people with hip or knee OA [39] in people with lumbar herniated disc or lumbar spinal stenosis [39] Construct validity: in people with hip or knee OA [39] in people with hip or knee OA [39] in people with lumbar herniated disc or lumbar spial stenosis for surgical decisions [117] Reliability: Internal consistency: no information in
						which populations [39,117] Test-retest reliability: no information in which populations [39,117]
Trust in Surgical Decision Scale	The level of patient trust that their surgeon will help them make a good decision about an operation [40]	 Knee OA (n=1) [40] Hip OA (n=1) [40] Lumbar herniated disc (n=1) [40] Lumbar spinal stenosis (n=1) [40] 	PROM	• 5 items [40]	 0-20 [40] Higher scores indicate higher trust [40] 	Validity: • Validated: - no information in which populations [40]
Decision Self Efficacy Scale	 Patient's level of confidence in various aspects of the decision making process [42]. Measures an individual's self-confidence or belief in their ability to make decisions and engage in shared decision making. (=the certainty an individual feels in making an informed choice) [41] Self-confidence in one's abilities to participate in shared-decision making [52] Measures self-confidence in one's abilities in decision making, including shared decision making [94] 	 Lumbar degenerative diseases (n=1) [41] Rheumatoid arthritis (n=1) [94] Osteoporosis (n=1) [42] Pain involving one or both knees (n=1) [52] 	PROM	• 11 items [41,42,52]	 0-100 [41] Scores of 0 and 100 indicate extremely low and extremely high self-efficacy, respectively [41] 	Validity: • Validated: - no information in which populations [42] • Discriminant validity: - in people with schizophrenia [52] • Convergent validity: - no information in which populations [41] • Correlation with other constructs: - no information in which populations [52] Reliability: • Reliable: - no information in which populations [42] • Internal consistency: - no information in which populations [42] (Internal consistency: - no information in which populations [41,52] (continued on next page)

Measure-ment instrument	Extracted construct	Condition(s) in which it was used	Type of measu- re	Number of items and subscales	Score, cut-off and interpretation	Information on measurement properties
Patient-Doctor Relation-ship Question- naire	 Patient's perception of their physician as effective and helpful [46] The relationship between the physician and the patient from the patient's perspective [54] 	 Knee OA (n=1) [46] Hip OA (n=1) [46] Lupus nephritis (n=1) [54] 	PROM	• 9 items [46,54]	 9–45 [46,54] Higher score indicating a greater patient's perception of the effectiveness and helpfulness of the physician [46]. Higher scores reflect a better relationship between patients and their doctors [54]. 	Validity: • Validated: - no information in which populations [46] Reliability: • Internal consistency: - in a primary care setting [54]
OPTION Scale	 Shared decision making [48] The extent to which clinicians sought to involve patients in decision making [76,78, 88,90,98] 	 Knee OA (n=1) [48] Rheumatoid arthritis (n=1) [88] Osteopenia or osteoporosis (n=3) [76, 78,90] Non-chronic low back pain (n=1) [98] 	CROM	• 5 items [88] • 12 items [48,98]	O-100 [48,76,88,98] Higher scores indicate higher levels of shared decision making [88]	Validity: • Validated: - no information in which populations [88,98] Reliability:
MASRI	• Medication adherence [54]	• Lupus nephritis (n=1) [54]	PROM	• 12 items [54]	• 0–100 [54]	Reliable: no information in which populations [88] Validity:
					which populations [54] • Predictive validity: - no information in which populations [54] Reliability: • Reliable:	
Beliefs about Medicines Question- naire	Patients' beliefs and concerns about taking medication for their condition [54,91]	 Rheumatoid arthritis (n=1) [91] Psoriatic arthritis (n=1) [91] Ankylosing spondylitis (n=1) [91] Lupus nephritis (n=1) [54] Osteoporosis (n=1) [93] 	PROM	 10 items [91] 18 items [54] 2 subscales (patient's beliefs about the necessity of medication and their concerns about it) [91] 	5-25 [91] Higher scores indicate stronger beliefs about the corresponding elements in each subscale translating into more negative beliefs about medicines [54]	 in people with systemic lupus erythematosus [54] Validity: Predictive validity: no information in which populations [54] Reliability: Internal consistency: no information in which populations [54,91] Test-retest reliability: no information in which populations
Interperso-nal Trust in a Physician	Overall patient trust in their individual physician [54]	• Lupus nephritis (n=1) [54]	PROM	• 10 items [54]	• Higher scores reflect higher levels of trust in the physician [54]	[54] Reliability: • Internal consistency: - no information in which populations [54] • Test-retest reliability:
Effective Consumer Scale	 Patients' perceptions of their ability to effectively manage and participate in their healthcare [79] The main skills and behaviors that people require to manage their health effectively [82] 	 Rheumatoid arthritis (n=1) [79] Osteoporosis (n=1) [82] 	PROM	• 17 items [79,82]	0-100 [79,82] A higher score indicates better disease management skills [79, 82]	 no information in which populations [54] Reliability: Test-retest reliability: no information in which populations [82] (continued on next page

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Measure-ment instrument	Extracted construct	Condition(s) in which it was used	Type of measu- re	Number of items and subscales	Score, cut-off and interpretation	Information on measurement properties
Medication Education Impact Question- naire	The impact of education intervention in addressing patient needs and facilitating shared decision making and self-management [80]	• Rheumatoid arthritis (n=1) [80]	PROM	29 items [80] 6 subscales (Information Quality, Active Communication, Coming to Terms with Diagnosis and Treatment, Self- management Role, Self- management Capacity, and Self-management Support) [80]	• Higher=better [80]	Reliability: Internal consistency: in people with rheumatic diseases [80] Test-retest reliability: in people with rheumatic diseases [80]
Morisky Medication Adherence Scale	• Self-reported adherence [81,91]	 Rheumatoid arthritis (n=2) [81,91] Psoriatic arthritis (n=2) [81,91] Ankylosing spondylitis (n=1) [91] 	PROM	• 4 items [81] • 8 items [91]	 0-4 [81] 0-8 [91] 0 indicating high adherence and 3-4 indicating low adherence [81] Higher scores representing more adherent behaviour [91] 	Reliability: • Internal consistency: - no information in which populations [91]
Perceived Involve-ment in Care Scale	Involvement in decision making [110]	• Acute low back pain (n=1) [110]	PROM	2 subscales (Doctor Facilitation subscale and Information Exchange subscale) [110]	0-100 [110] Higher scores indicating a greater involvement of the health care provider (doctor facilitation subscale) [110] Higher scores indicating a greater extend of active information seeking (information exchange subscale) [110]	Reliability: • Internal consistency: - no information in which populations [110]
Satisfaction with Information about Medicines Scale	 Satisfaction with the amount of information received [91] Patient satisfaction [93] 	 Rheumatoid arthritis (n=1) [91] Psoriatic arthritis (n=1) [91] Ankylosing spondylitis (n=1) [91] Osteoporosis (n=1) [93] 	PROM	• 21 items [91]	 0-21 [91] Higher scores indicating a higher degree of overall satisfaction with the amount of information received [91] 	Reliability: • Internal consistency: - no information in which populations [91]
Decision Evaluation Scales	• Assesses (1) informed choice: the patient's perception of the quality of the received information; (2) decision control: the patient's perceived level of control over the decision in terms of feelings of regret, anxiety and deciding under pressure; and (3) satisfaction-uncertainty: the extent to which a patient is satisfied or still has doubts about the decision [91].	 Rheumatoid arthritis (n=1) [91] Psoriatic arthritis (n=1) [91] Ankylosing spondylitis (n=1) [91] 	PROM	15 items [91] 3 subscales (informed choice, decision control and satisfaction uncertainty) [91]	3–15 [91] Higher scores indicating higher levels of informed choice, decision control and higher satisfaction (less uncertainty) [91].	Reliability: • Internal consistency: - no information in which populations [91]
Cologne Patient Question- naire	Trust in the physician and need for information [91]	 Rheumatoid arthritis (n=1) [91] Psoriatic arthritis (n=1) [91] Ankylosing spondylitis (n=1) [91] 	PROM	7 items [91]2 subscales (trust in the physician and need for information) [91]	 Higher score indicating greater trust [91] Higher score indicating higher need for information [91] 	Reliability: Internal consistency: no information in which populations
Decision readiness	• Decision readiness [119]	(n=1) [91] • Knee OA (n=1) [119]	PROM	• 1 item [119]	Higher answers (like "very") indicate greater decision readiness [119]	[91] Validity: • Validated: - no information in which populations [119]
Stage of Decision- Making Scale	• Stage of decision making [25,122]	• Knee OA (n=2) [25, 122] • Hip OA (n=2) [25,122]	PROM	• 1 item [25]		Validity: • Validated: (continued on next page)

Measure-ment instrument	Extracted construct	Condition(s) in which it was used	Type of measu- re	Number of items and subscales	Score, cut-off and interpretation	Information on measurement properties
Treatment intention	• Treatment intention [123]	• Subacromial pain syndrome (n=1) [123]	PROM	• 1 item [123]	0–100 [123] Higher scores indicate higher intention to try surgery [123]	- no information in which populations [122] Reliability: • Test-retest reliability: - no information in which populations [123] Responsiveness:
Satisfaction with decision and decision making process	Satisfaction with decision and decision making process [91]	 Rheumatoid arthritis (n=1) [91] Psoriatic arthritis (n=1) [91] Ankylosing spondylitis (n=1) [91] 	PROM	6 scales (satisfaction with participation, satisfaction with amount of received information, informed choice, decision control, satisfaction-uncertainty, and consistency with personal values) [91]	1–5 for satisfaction with participation subscale [91] Higher score indicating higher levels of satisfaction with participation (for satisfaction with participation subscale)	Sensitive to change [123] Reliability Internal consistency: within the Nota's study sample [91]
Decision process Knowledge on	 Decision process [102] Patient knowledge on 	 Knee OA (n=1) [102] Hip OA (n=1) [102] Acute low back pain 	PROM	• 4 items [102]	[91] • 0-100 [102] • Higher score indicating more involvement [102]	Reliability Internal consistency: within the Sepucha's study sample [102] Test-retest reliability: within the Sepucha's study sample [102] Reliability:
acute low back pain	acute low back pain [110]	(n=1) [110]				Internal consistency: within the Simon's study sample [110]

OA: osteoarthritis

PROM: patient reported outcome measure

DQI: Decision Quality Instruments

FAPI: Fragebogen zur Arzt-Patienten-Interaktion

OPTION Scale: Observing Patient Involvement in Decision Making instrument

CROM: clinician reported outcome measure.

MASRI: Medication Adherence Self-Report Inventory

Appendix C. Characteristics of the measurement instruments without any measurement properties reported by the included studies

of 7 out of 8 correct answers onsidered adequate rheumatoid is knowledge
al number of items answered ly
[34,64,81]
scores indicating more patient
ion [34,64,81]

(continued on next page)

Measured construct and its definition	Condition(s) in which it was used	Type of measure	Number of items and subscales	Score, cut-off, and interpretation
(nowledge of rheumatoid arthritis and treatment, and perceived self-management behaviors (Partners in Health Scale)	Rheumatoid arthritis [80]	PROM	11 items	0–88 [80]Higher score indicating better knowledge [80]
Knowledge about etanercept	Rheumatoid arthritis [87]	PROM	12 items	0–12
Patient knowledge Knowledge about neuromuscular scoliosis treatment	Osteoporosis [90] Parent(s) of a child with	PROM	5 items	
Knowledge about hip or knee OA disease progression and	neuromuscular scoliosis [108] • Hip OA [109]	PROM	5 items	0–5
total hip or knee arthroplasty*	• Knee OA [109]			
Knowledge about OA and total knee replacement	Knee OA [115]	PROM	7 items 4 items	0–100
Knowledge about treatment options and risks	Hip OA [118]Knee OA [118]	PROM	4 Items	0–4
Knowledge about options*	Subacromial pain syndrome [123]	PROM	7 items	0–7
TAGE OF DECISION	V700 OA [44]		1 itom	
Preparation to make a decision on their preference* Decision making stage*	Knee OA [44] • Hip OA [63]	PROM	1 item 1 item	
	• Knee OA [63]			
Stage of decision making*	Anterior shoulder dislocation [64]	PROM	1 item	
Stage of decision making*	Spinal stenosis [70]	PROM	1 item	
Stage of decision making	Hip OA [109]Knee OA [109]	PROM	1 item	
Stage of decision making	• Knee OA [109] • Hip OA [118]			
от от то т	• Knee OA [118]			
Stage of decision making*	Knee OA [120]	PROM	1 item	1–6
VALUES AND PREFERENCES				
Value concordance	Knee OA [27]	Analysis	1 :+	
Treatment preference* Values	Knee OA [27]	PROM PROM	1 item 2 items	
Choice predisposition	Osteoporosis [42] Osteoporosis [42]	PROM	1 item	
Freatment alignment	Knee OA [48]	Analysis	1 Itciii	
Values	Knee OA [48]	PROM	1 item	
/alues	Rheumatoid arthritis [51]	PROM	10 items	
Preference towards biosimilars*	Rheumatoid arthritis [53]	PROM	1 item	• 0–10
	 Psoriatic arthritis [53] Granulomatosis with polyangiitis [53] Juvenile idiopathic arthritis [53] Other rheumatic diseases [53] 			preferences for biosimilars
nfluential values of patient regarding drug treatment	Rheumatic diseases [57]	Analysis		
Treatment preference*	Hip OA [63]Knee OA [63]	PROM	1 item	
Freatment alignment with evidence-based treatment	Anterior shoulder dislocation [64]	Analysis		
mportance*	Knee OA [66]	PROM	3 items	
/alues and goals*	Knee OA [66]	PROM	2 items	
Summarizing pros and cons*	Knee OA [66]	PROM	2 items	
Treatment concordance*	Knee OA [68]	PROM	1 item	
Freatment preference fit index	Rheumatoid arthritis [83]	Analysis	1 item	
Treatment preference Treatment preference	Knee OA [97] • Hip OA [109]	PROM PROM	1 item	
realism preservice	• Knee OA [109]	1 1(01)1	1 Item	
/alues*	Psoriatic arthritis [114]	Interview	3 questions	
mportance given to certain characteristics and potential	Knee OA [115]	PROM		
outcomes of OA treatments Creatment preference	Vnee OA [115]	PROM	1 item	
Treatment preference	Knee OA [115] • Hip OA [118]	PROM	1 item	
realism profession	• Knee OA [118]	11(01)1		
DECISION				
Treatment decision	Knee OA [27]	Chart		
reatment choice*	Hip OA [36]	review PROM	1 item	
readment choice	Knee OA [36]	1 IOIVI	1 ICIII	
nformed choice	Hip OA [39] Knee OA [39] Lumbar herniated disc [39]	Analysis		
	Lumbar spinal stenosis [39]			
Choice	Osteoporosis [42]	PROM	1 item	
nformed choice	Rheumatoid arthritis [51]	Analysis		
nformed choice Confidence with the decision (Subscale of the Combined	Rheumatoid arthritis [50] Rheumatoid arthritis [50]	Analysis PROM	10 items	
Outcome Measure for Risk Communication) Freatment choice	Upper-extremity conditions [73]	PROM	1 item	
Treatment choice	Displaced diaphyseal clavicle fractures			
	[77]			(
				(continued on next

Measured construct and its definition	Condition(s) in which it was used	Type of measure	Number of items and subscales	Score, cut-off, and interpretation
Decision to start biphosphonates	Osteoporosis [78]			
Decision regret*	• Hip OA [102]	PROM	1 item	
recision regret	• Knee OA [102]	TROM	1 Item	
Decision confidence*	• Hip OA [102]	PROM	1 item	
ecision confidence	=	PROM	1 Item	
Wanter out wanter d	• Knee OA [102]	Doored		
reatment received	• Hip OA [106]	Record		
	• Knee OA [106]			
nformed, patient-centred decision	• Hip OA [106]	Analysis		
	• Knee OA [106]			
reatment decision	Osteopenia or osteoporosis [111]	PROM		
Pecision quality*	• Hip OA [113]	Analysis		
	• Knee OA [112,113]			
nformed, patient-centred decision	• Hip OA [117]	Analysis		
	• Knee OA [117]			
	 Lumbar herniated disc [117] 			
	 Lumbar spinal stenosis [117] 			
atient had made their definitive decision after the first visit	 Hip OA [118] 	PROM		
	• Knee OA [118]			
Decision readiness*	Knee OA [120]	PROM	1 item	
reatment choice	Trapeziometacarpal arthritis [121]	•	-	
reatment choice	Hip OA[122]			
	Knee OA [122]			
nformed, patient centred decision	• Hip OA [122]	Analysis		
mormed, patient centred decision	• Knee OA [122]	1 11101 y 313		
nformed choice	Knee OA [122] Subacromial pain syndrome [123]	Apolyeis		
	Subacromiai pain syndrome [123]	Analysis		
VILLINGNESS	Warr OA FORT			
Villingness to have surgery	Knee OA [27]			
Villingness to have acupuncture*	Back pain [32]	PROM	1 item	
Villingness to take a (new) biological treatment* (Choice	Rheumatoid arthritis [51]	PROM	1 item	
Predisposition Scale)				
Villingness to change to a biosimilar*	 Rheumatoid arthritis [53] 	PROM	1 item	
	 Ankylosing spondylitis [53] 			
	 Psoriatic arthritis [53] 			
	 Granulomatosis with polyangiitis 			
	[53]			
	Juvenile idiopathic arthritis [53]			
	Other rheumatic diseases [53]			
Villingness to have surgery (Choice Predisposition Scale)	Knee OA [59]	PROM	1 item	
Villingness to undergo total knee replacement if	Knee OA [65]	PROM	1 item	
	Rice on [00]	TROM	1 Item	
recommended by the surgeon Villingness to consider total knee replacement*	Vnoc OA [120]	PROM	1 item	
DURATION OF ENCOUNTER	Knee OA [120]	PROM	1 Item	
	YY' OA FOCT			
ength of consultation time	• Hip OA [36]	Analysis		
	• Knee OA [36]			
Ouration of the visit	Orthopaedic surgery [37]	Analysis		
Ouration of consultations	Knee OA [68]	Analysis		
Ouration of encounters	Osteopenia or osteoporosis [78]	Analysis		
ATISFACTION				
atisfaction with the visit*	 Hip OA [36] 	PROM	1 item	0–10
	 Knee OA [36] 			
Overall satisfaction*	 Hip OA [39] 	PROM	1 item	
	• Knee OA [39]			
	Lumbar herniated disc [39]			
	Lumbar spinal stenosis [39]			
atisfaction with decision	Acute musculoskeletal pain [60]	PROM	1 item	
atisfaction with treatment	•	PROM	1 item	
	Acute musculoskeletal pain[60]			
atisfaction with treatment choice*	Orthopaedic injuries[67]	PROM	1 item	
atisfaction with consultation	Knee OA[68]	PROM	1 item	
atisfaction*	Dupuytren contracture[69]	Interview	2 questions	
atisfaction with care*	Rheumatoid arthritis[72]	PROM	1 item	
	Ankylosing spondylitis[72]			
atisfaction with the visit	Upper-extremity conditions[73]	PROM	1 item	
Overall satisfaction with care*	Spinal disorders[74]	PROM	1 item	
Verali satisfaction with the	Musculoskeletal pain[75]	PROM	18 items	0–100
atisfaction with pain care (Pain Treatment Satisfaction				Higher scores indicate greater
				satisfaction
atisfaction with pain care (Pain Treatment Satisfaction				
atisfaction with pain care (Pain Treatment Satisfaction Scale)	Rheumatoid arthritis[83]	PROM	1 item	
atisfaction with pain care (Pain Treatment Satisfaction Scale) atisfaction with treatment*	Rheumatoid arthritis[83] Knee OA[86]	PROM PROM	1 item 3 items	
atisfaction with pain care (Pain Treatment Satisfaction Scale) atisfaction with treatment* atisfaction with results of knee replacement*	Knee OA[86]	PROM PROM	1 item 3 items	
atisfaction with pain care (Pain Treatment Satisfaction Scale) atisfaction with treatment* atisfaction with results of knee replacement* atisfaction with knowledge transfer	Knee OA[86] Osteoporosis[90]	PROM	3 items	
atisfaction with pain care (Pain Treatment Satisfaction Scale) atisfaction with treatment* atisfaction with results of knee replacement* atisfaction with knowledge transfer	Knee OA[86] Osteoporosis[90] • Rheumatoid arthritis[92]			
atisfaction with pain care (Pain Treatment Satisfaction	Knee OA[86] Osteoporosis[90] Rheumatoid arthritis[92] Psoriatic arthritis[92]	PROM	3 items	
atisfaction with pain care (Pain Treatment Satisfaction Scale) atisfaction with treatment* atisfaction with results of knee replacement* atisfaction with knowledge transfer atisfaction with decision making process*	Knee OA[86] Osteoporosis[90] Rheumatoid arthritis[92] Psoriatic arthritis[92] Ankylosing spondylitis[92]	PROM	3 items	
atisfaction with pain care (Pain Treatment Satisfaction Scale) atisfaction with treatment* atisfaction with results of knee replacement* atisfaction with knowledge transfer	Knee OA[86] Osteoporosis[90] Rheumatoid arthritis[92] Psoriatic arthritis[92]	PROM	3 items	

Measured construct and its definition	Condition(s) in which it was used	Type of measure	Number of items and subscales	Score, cut-off, and interpretation
Satisfaction with decision making process Patient's satisfaction with how their treatment turned out	Knee OA[97] • Hip OA[103] • Knee OA[103] • Lumbar herniated disc[103]	PROM	12 items	
Patient's satisfaction with their current pain and symptoms	Lumbar spinal stenosis[103] Hip OA[103] Knee OA[103] Lumbar herniated disc[103] Lumbar herniated disc[103]			
Satisfaction with treatment*	Lumbar spinal stenosis[103]Hip OA[106]Knee OA[106]	PROM	1 item	
Satisfaction with their current pain	Hip OA[106]Knee OA[106]	PROM	1 item	
Satisfaction with shared decision making	Rheumatoid arthritis[107]Psoriatic arthritis[107]Axial spondylarthritis[107]			
Parent's satisfaction with shared decision making Satisfaction regarding education and knowledge*	Neuromuscular scoliosis[108] • Hip OA[109] • Knee OA[109]	PROM PROM	1 item	
Satisfaction with received information* Satisfaction with decision making process	Psoriatic arthritis[114] Knee OA[115]	Interview PROM	2 questions 12 items	0–100
Overall satisfaction with care* Patients' satisfaction with the given information, the clinic,	Midshaft clavicle fracture [116] • Hip OA[118] • Knee OA[118]	PROM PROM	7 items 3 items	
and the physician Satisfaction with the visit* Overall treatment satisfaction* EXPECTATIONS	Trapeziometacarpal arthritis[121] Trapeziometacarpal arthritis[121]	PROM PROM	1 item 1 item	
Realistic expectations*	Osteoporosis[42]	PROM	5 items	The score for realistic expectations was the percent of accurate responses
Expectations about knee replacement post-surgery* Realistic expectations*	Knee OA[86] • Hip OA[113] • Knee OA[113]	PROM Analysis	1 item	out of the five questions.
RISK Risk communication (Subscale of the Combined Outcome Measure for Risk Communication)	Rheumatoid arthritis[50]	PROM	10 items	
Perceptions of cognitive and affective risk*	 Rheumatoid arthritis[53] Ankylosing spondylitis[53] Psoriatic arthritis[53] Granulomatosis with polyangiitis [53] Juvenile idiopathic arthritis[53] Other rheumatic diseases[53] 	PROM	1 item	
SUPPORT/RELATIONSHIP Practical and emotional support received by accompanied patients during the decision process*	 Rheumatoid arthritis[53] Ankylosing spondylitis[53] Psoriatic arthritis[53] Granulomatosis with polyangiitis [53] Juvenile idiopathic arthritis[53] Other rheumatic diseases[53] 	PROM	2 items	
Support* Relationship and communication with clinicians*	Dupuytren contracture[69] Psoriatic arthritis[114]	PROM Interview	1 item 3 questions	
OTHER Explanation understanding*	 Rheumatoid arthritis[53] Ankylosing spondylitis[53] Psoriatic arthritis[53] Granulomatosis with polyangiitis [53] 	PROM	1 item	
Reassurance*	 Juvenile idiopathic arthritis[53] Other rheumatic diseases[53] Rheumatoid arthritis[53] Ankylosing spondylitis[53] Psoriatic arthritis[53] Granulomatosis with polyangiitis [53] 	PROM	1 item	
Preference in receiving information accompanied*	 Juvenile idiopathic arthritis[53] Other rheumatic diseases[53] Rheumatoid arthritis[53] Ankylosing spondylitis[53] Psoriatic arthritis[53] Granulomatosis with polyangiitis 	PROM	1 item	
	[53]Juvenile idiopathic arthritis[53]			(continued on next page)

Measured construct and its definition	Condition(s) in which it was used	Type of measure	Number of items and subscales	Score, cut-off, and interpretation
	Other rheumatic diseases[53]			
Awareness of making a preference-sensitive decision	Anterior shoulder dislocation[64]			
Discussion of knee pain with primary care provider*	Knee OA[66]	Interview	7 questions	
Readiness*	Knee OA[66]	PROM	2 items	
Confidence*	Knee OA[66]	PROM	1 item	
Received information*	Rheumatoid arthritis[72]	PROM	3 items	
teerved information	Ankylosing spondylitis[72]	1110111	o items	
Unmet health care needs*	Rheumatoid arthritis[72]	PROM	2 items	
	 Ankylosing spondylitis[72] 			
Overall experience with decision making process*	Knee OA[96]	PROM	2 items	
Confidence with current understanding*	Psoriatic arthritis[114]	Interview	2 questions	
Beliefs about effectiveness of joint replacement*	Knee OA[120]	PROM	1 item	
Change of treatment	Trapeziometacarpal arthritis[121]	Analysis	1 110111	
Perception of the physician's empathic understanding during	Trapeziometacarpal arthritis[121]	PROM	10 items	0-50[121]
the office visit (Consultation and Relational Empathy Scale)	Taponomentupa manta(121)	11.0.11	10 1101110	Higher score indicating greater empathy[121]
Attitudes towards surgery	Subacromial pain syndrome[123]	PROM	3 items	3–21
SHARED DECISION MAKING				
nformed decision making*	Orthopaedic surgery[37]	CROM	9 elements	
Shared decision making process*	Anterior cruciate ligament injury[56]	PROM	10 items	
Specific aspects of shared decision making*	Rheumatoid arthritis[61]	CROM		
Components of shared decision making*	Orthopaedic injuries[67]	PROM	2 items	
Decision making process*	Dupuytren contracture[69]	Interview	11 questions	
Involvement in decisions*	Rheumatoid arthritis[72]	PROM	2 items	
HAOLACHICHT III (ICC1910119		r KOIVI	∠ 1(CHI)	
Shared decision making*	 Ankylosing spondylitis[72] Lumbar spine conditions (postsurgery) [74] 	PROM	4 items	
Medication-related shared decision making (Consumer Assessment of Healthcare Provider and System survey)	Musculoskeletal pain[75]		3 items	
Preferences for shared decision making*	Rheumatoid arthritis[83]	PROM	2 items	
Patients' experience of shared decision making	Anterior cruciate ligament injury[84]	Interview		
Patient-reported shared decision making*	Non-chronic low back pain[98]	PROM	1 item	
The reasons women present when expressing hesitation about initiation of bisphosphonates during primary care consultations with clinicians and how these clinicians	Osteoporosis[99]	CROM	7 categories	
react*				
Decision making participation	• Hip OA[109]	PROM	2 items	
	• Knee OA[109]			
Preference for participation	Acute low back pain[110]	PROM	1 item	
Patient-reported shared decision making	Osteopenia or osteoporosis[111]	PROM		
Control when making a decision*	Psoriatic arthritis[114]	Interview	1 question	
Decision making process*	Psoriatic arthritis[114]	Interview	3 questions	
ADHERENCE		-		
Compliance	Juvenile idiopathic arthritis[47]	Analysis		
Persistence	Juvenile idiopathic arthritis[47]	Analysis		
Medication adherence (assumed as compliance)	Osteoporosis [49]	Analysis		
Medication adherence (assumed as persistence)	Osteoporosis [49]	Analysis		
	•	viigi 3918		
Continuance rate of treatment	Rheumatic diseases[57]	Medical		
Having undergone arthroplasty	• Hip OA[62]			
	 Knee OA[62] 	record		
	TT OATCET			
Receipt of total knee replacement	Knee OA[65]	Medical		
		record		
Total knee replacement rates	Knee OA[68]	record		
Cotal knee replacement rates Analgesic adherence (Pain Medication in Primary Care			4 items	
Cotal knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire)	Knee OA[68] Musculoskeletal pain[75]	record PROM	4 items	
Cotal knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire) Primary adherence	Knee OA[68] Musculoskeletal pain[75] Osteoporosis[78]	record PROM Analysis	4 items	
Cotal knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire) Primary adherence Secondary adherence	Knee OA[68] Musculoskeletal pain[75]	PROM Analysis Analysis	4 items	
Cotal knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire) Primary adherence Secondary adherence	Knee OA[68] Musculoskeletal pain[75] Osteoporosis[78]	record PROM Analysis	4 items	
Cotal knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire) Primary adherence decondary adherence Medication adherence*	Knee OA[68] Musculoskeletal pain[75] Osteoporosis[78] Osteoporosis[78]	PROM Analysis Analysis		
Receipt of total knee replacement Fotal knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire) Primary adherence Secondary adherence Medication adherence* Medication adherence Persistence	Knee OA[68] Musculoskeletal pain[75] Osteoporosis[78] Osteoporosis[78] Osteoporosis[90]	PROM Analysis Analysis PROM		
Total knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire) Primary adherence Secondary adherence Medication adherence* Medication adherence	Knee OA[68] Musculoskeletal pain[75] Osteoporosis[78] Osteoporosis[90] Osteoporosis[90]	PROM Analysis Analysis PROM Analysis		
Cotal knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire) Primary adherence Secondary adherence Medication adherence* Medication adherence Persistence Densistence Compliance	Knee OA[68] Musculoskeletal pain[75] Osteoporosis[78] Osteoporosis[90] Osteoporosis[90] Osteoporosis[90]	PROM Analysis Analysis PROM Analysis Analysis Analysis		
Cotal knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire) Primary adherence Secondary adherence Medication adherence* Medication adherence Persistence Compliance Attendance at routine clinics and self-report compliance	Knee OA[68] Musculoskeletal pain[75] Osteoporosis[78] Osteoporosis[90] Osteoporosis[90] Osteoporosis[90] Osteoporosis[90] Osteoporosis[93]	PROM Analysis Analysis PROM Analysis Analysis Analysis		
Cotal knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire) Primary adherence Primary adherence Medication adherence* Medication adherence Persistence Compliance Attendance at routine clinics and self-report compliance (Medication Adherence Report Scale)	Knee OA[68] Musculoskeletal pain[75] Osteoporosis[78] Osteoporosis[90] Osteoporosis[90] Osteoporosis[90] Osteoporosis[93] Osteoporosis[93]	PROM Analysis Analysis PROM Analysis Analysis Analysis		
Cotal knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire) Primary adherence Primary adherence Medication adherence* Medication adherence Persistence Compliance Attendance at routine clinics and self-report compliance (Medication Adherence Report Scale) Having undergone total knee replacement	Knee OA[68] Musculoskeletal pain[75] Osteoporosis[78] Osteoporosis[90] Osteoporosis[90] Osteoporosis[90] Osteoporosis[93] Osteoporosis[93] Knee OA[97]	PROM Analysis Analysis PROM Analysis Analysis Analysis		
Total knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire) Primary adherence Primary adherence Primary adherence Medication adherence* Medication adherence Persistence Compliance Attendance at routine clinics and self-report compliance (Medication Adherence Report Scale) Having undergone total knee replacement	Knee OA[68] Musculoskeletal pain[75] Osteoporosis[78] Osteoporosis[90] Osteoporosis[90] Osteoporosis[90] Osteoporosis[93] Osteoporosis[93] Osteoporosis[93] Knee OA[97] • Hip OA[100]	PROM Analysis Analysis PROM Analysis Analysis Analysis Analysis		
Cotal knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire) Primary adherence Secondary adherence Medication adherence* Medication adherence Persistence Compliance Attendance at routine clinics and self-report compliance	Knee OA[68] Musculoskeletal pain[75] Osteoporosis[78] Osteoporosis[90] Osteoporosis[90] Osteoporosis[90] Osteoporosis[93] Osteoporosis[93] Knee OA[97] • Hip OA[100] • Knee OA[100]	PROM Analysis Analysis PROM Analysis Analysis Analysis Analysis		
Cotal knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire) Primary adherence Primary adherence Medication adherence* Medication adherence Persistence Compliance Attendance at routine clinics and self-report compliance (Medication Adherence Report Scale) Having undergone total knee replacement	Knee OA[68] Musculoskeletal pain[75] Osteoporosis[78] Osteoporosis[90] Osteoporosis[90] Osteoporosis[90] Osteoporosis[93] Osteoporosis[93] Knee OA[97] • Hip OA[100] • Knee OA[100]	PROM Analysis Analysis PROM Analysis Analysis Analysis Analysis		
Total knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire) Primary adherence Secondary adherence Medication adherence* Medication adherence Persistence Compliance Attendance at routine clinics and self-report compliance (Medication Adherence Report Scale) Having undergone total knee replacement Having undergone surgery	Knee OA[68] Musculoskeletal pain[75] Osteoporosis[78] Osteoporosis[90] Osteoporosis[90] Osteoporosis[90] Osteoporosis[93] Osteoporosis[93] Knee OA[97] • Hip OA[100] • Knee OA[100] • Lumbar herniated disc[100] • Lumbar spinal stenosis[100]	PROM Analysis Analysis PROM Analysis Analysis Analysis Medical record		
Total knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire) Primary adherence Primary adherence Primary adherence Medication adherence* Medication adherence Persistence Compliance Attendance at routine clinics and self-report compliance (Medication Adherence Report Scale) Having undergone total knee replacement	Knee OA[68] Musculoskeletal pain[75] Osteoporosis[78] Osteoporosis[90] Osteoporosis[90] Osteoporosis[90] Osteoporosis[93] Osteoporosis[93] Knee OA[97] • Hip OA[100] • Lumbar herniated disc[100] • Lumbar spinal stenosis[100] • Hip OA[103]	PROM Analysis Analysis PROM Analysis Analysis Analysis Analysis Medical record		
Total knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire) Primary adherence Secondary adherence Medication adherence* Medication adherence Persistence Compliance Attendance at routine clinics and self-report compliance (Medication Adherence Report Scale) Having undergone total knee replacement Having undergone surgery	Knee OA[68] Musculoskeletal pain[75] Osteoporosis[78] Osteoporosis[90] Osteoporosis[90] Osteoporosis[90] Osteoporosis[93] Osteoporosis[93] Knee OA[97] • Hip OA[100] • Knee OA[100] • Lumbar herniated disc[100] • Limbar spinal stenosis[100] • Hip OA[103]	PROM Analysis Analysis PROM Analysis Analysis Analysis Medical record		
Cotal knee replacement rates Analgesic adherence (Pain Medication in Primary Care Patient Questionnaire) Primary adherence Secondary adherence Medication adherence Medication adherence Persistence Compliance Attendance at routine clinics and self-report compliance (Medication Adherence Report Scale) Having undergone total knee replacement Having undergone surgery	Knee OA[68] Musculoskeletal pain[75] Osteoporosis[78] Osteoporosis[90] Osteoporosis[90] Osteoporosis[90] Osteoporosis[93] Osteoporosis[93] Osteoporosis[93] Knee OA[97] • Hip OA[100] • Knee OA[100] • Lumbar herniated disc[100] • Lumbar spinal stenosis[100] • Hip OA[103] • Knee OA[103]	PROM Analysis Analysis PROM Analysis Analysis Analysis Analysis Medical record		
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Measured construct and its definition	Condition(s) in which it was used	Type of measure	Number of items and subscales	Score, cut-off, and interpretation	
Having undergone total knee replacement	• Knee OA[113] Knee OA[115]	PROM	1 item		

PROM: patient reported outcome measure

OA: osteoarthritis

CROM: clinician reported outcome measure.

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