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Barriers and enablers in primary care clinicians’ management of osteoarthritis: protocol for a systematic review and qualitative evidence synthesis

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ABSTRACT

Introduction: Osteoarthritis is a highly prevalent and disabling condition. Primary care management of osteoarthritis is generally suboptimal despite evidence for several modestly effective interventions and the availability of high-quality clinical practice guidelines. This report describes a planned study to synthesise the views of primary care clinicians on the barriers and enablers to following recommended management of osteoarthritis, with the aim of providing new interpretations that may facilitate the uptake of recommended treatments, and in turn improve patient care.

Methods and analysis: A systematic review and meta-synthesis of qualitative studies. 5 databases will be searched using key search terms for qualitative research, evidence-based practice, clinical practice guidelines, osteoarthritis, beliefs, perceptions, barriers, enablers and adherence. A priori inclusion/exclusion criteria include availability of data from primary care clinicians, reports on views regarding management of osteoarthritis, and studies using qualitative methods for both data collection and analysis. At least 2 independent reviewers will identify eligible reports, conduct a critical appraisal of study conduct, extract data and synthesise reported findings and interpretations. Synthesis will follow thematic analysis within a grounded theory framework of inductive coding and iterative theme identification. The reviewers plus co-authors will contribute to the meta-synthesis to find new themes and theories. The Confidence in the Evidence from Reviews of Qualitative research (CERQual) approach will be used to determine a confidence profile of each finding from the meta-synthesis. The protocol has been registered on PROSPERO and is reported using the Preferred Reporting Items for Systematic Review and Meta-Analyses Protocols (PRISMA-P) guidelines.

Ethics and dissemination: Ethical approval is not required. The systematic review will be published in a peer-reviewed journal. The results will help to inform policy and practice and assist in the optimisation of management for people with osteoarthritis.

PROSPERO registration number: CRD42015027543.

Strengths and limitations of this study

- The search strategy is designed to be comprehensive and informed by Cochrane review methods, published recommendations for optimal qualitative research identification, and a previously published protocol.
- Inclusion/exclusion criteria and data extraction have been determined a priori to reduce bias in sourcing data, and study screening, data extraction, critical appraisal of study conduct and data analysis will be independently conducted by more than one reviewer with a further reviewer available for arbitration to reach consensus.
- The first limitation is removal of data from the original contexts and the participant quotes are no longer linked to the context of the original questions or the participant’s setting.
- The second limitation is that the synthesis relies on the data presented in each of the included reports which may not reflect the full analysis of the original data.
- The inclusion of only English language publications means there is potential for cultural and publication bias in the findings.

INTRODUCTION

Osteoarthritis (OA) is a major global public health problem causing significant pain and disability, and is now ranked 13th in global causes of years lived with disability.2 There is no cure, but an extensive body of research has provided evidence to support use of a range of modestly effective treatments for symptom and function management.3–6 Evidence-based and expert consensus-based clinical practice guidelines (CPGs) have been produced to provide recommendations for effective treatments and best practice for OA management.7–13 Across these CPGs, conservative non-drug non-surgical care is advocated as the cornerstone for management of OA in all joints. This care includes...
education about the disease process, pain mechanisms and treatment options, and promotion of self-management with emphasis on positive behavioural changes, in particular exercise for all patients and weight loss for overweight or obese patients, regardless of joint(s) affected. Pharmacological options most recommended include acetylsalicylic acid (aspirin) or paracetamol as first-line, and non-steroidal anti-inflammatory drugs (topical or oral) as second-line options. Guidelines for management of knee OA uniformly advise against the use of arthroscopic debridement and/or lavage. Total knee or hip arthroplasty is recommended when people have severe symptomatic knee or hip OA.

While there is a large amount of agreement and overlap across recommendations in CPGs, some inconsistencies are apparent. Inconsistencies may occur because of the specific focus of the guideline, because of variations in quality and rigour of guideline development procedures, and because of changes in evidence over time. For example, recommendations have been specifically produced for management of hand OA, while others include all OA. Guidelines have also been produced which focus only on non-surgical or non-pharmacological management of OA. Quality assessment using the Appraisal of Guidelines Research and Evaluation (AGREE) Instrument indicates suboptimal quality of many CPGs for OA.

Guidelines use a range of evidence to formulate recommendations with high-quality evidence used where it exists but where it is lacking, recommendations may be based on lower quality evidence and/or expert opinion. Despite a lack of high-quality evidence, walking aids and thermal modalities are widely endorsed by CPGs for OA, while acupuncture, knee braces, heel wedges, intra-articular hyaluronans, glucosamine and chondroitin remain controversial. With time, evidence for or against some existing recommendations strengthens and evidence to support new options emerges. For example, the recommendation around the use of imaging for OA diagnosis has evolved, with older guidelines often recommending plain film X-ray to assist in the diagnosis of OA, while the recent National Institute for Health and Care Excellence (NICE) guidelines recommend that OA should be diagnosed clinically and without imaging.

OA is mostly diagnosed and managed in primary care settings and mostly by general practitioners (GPs; ie, family doctors). Some conservative interventions with recommendations for their use in CPGs, in particular exercise, weight loss, pain management advice and provision of other joint support or protection devices, may also be provided in primary care by allied health practitioners including physiotherapists, occupational therapists, exercise physiologists, podiatrists, pharmacists, practice nurses, dieticians and nutritionists. Survey and questionnaire data have shown that care received by individuals with OA in primary care settings is often inconsistent with broad CPG recommendations. Suboptimal care has been demonstrated across a number of quality domains including provision of effective treatments, safety, access to educational material and support for self-management.

For example, the Australian CareTrack study showed that only 43% of people with OA received recommended care. In particular, non-drug, non-surgical interventions are not given the importance by GPs that is recommended by all CPGs, while prescribing patterns appear to be better aligned with recommendations for pharmacological treatments. However, one study found there may be higher levels of prescription of more potent opioids than evidence suggests is warranted.

The development and dissemination of CPGs has been suggested as one method for improving the alignment of practice with evidence for effectiveness. However, previous research has highlighted that the availability of CPGs does not necessarily lead to evidence-based practice. Variations in accessing and uptake of research evidence and CPGs occur between different clinicians, between different sources of evidence and between different recommendations within guidelines. Clinician variations occur for many reasons, from differences in preferences for accessing information to strength of personal beliefs about health interventions. How concrete a recommendation is written, how complex or difficult the procedure is to provide, how credible the recommendation seems, and how feasible it is to implement locally will also influence uptake. These challenges are experienced across many health disorders where CPGs have been produced. With or without CPGs, change to more effective practice is often slow and inconsistent.

Qualitative methodologies are appropriate for exploring the nature of perceptions, beliefs, barriers and enablers that can influence whether practices align to evidence and recommendations. Qualitative synthesis of primary qualitative studies can pull together findings from across different settings and generate new theoretical or conceptual models. We have planned a qualitative synthesis exploring the barriers, enablers and/or beliefs and perceptions that may act as barriers or enablers to implementation of effective treatments and/or CPG recommendations for OA within the context of primary care practice. The findings may be helpful in informing our understanding of the complexity of implementing evidence-based guidelines for OA management and lead to innovations in addressing the evidence practice gap.

**METHODS**

The study has been registered on PROSPERO (http://www.crd.york.ac.uk/PROSPERO; 4/11/2015, registration number CRD42015027543). The reporting of this protocol is in accordance with the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) Statement for Protocols (PRISMA-P checklist) (see...
Eligibility criteria

Articles that report empirical data from primary care clinicians who provide treatments for OA (GPs or family doctors, physiotherapists, occupational therapists, exercise physiologists, podiatrists, pharmacists, practice nurses, dieticians and nutritionists) will be included. Views on barriers, enablers and/or perceptions regarding management of OA that were based on OA CPG recommendations will be sought. Studies will be excluded if they did not utilise qualitative methods for data collection and analysis. Mixed-methods studies will be included if the qualitative data are reported separately. Studies with multiple participant cohorts will be included if eligible primary care clinician groups’ data are reported separately. Articles reporting on other types of arthritis will be included only if data on OA are reported separately. Only English language reports will be included.

Identification and selection of studies

The search strategy was based on the one published for MEDLINE by Slade et al\(^3\) and adapted for OA and other databases. The strategy aims to identify all relevant articles published in peer-reviewed journals, and thus be a comprehensive assembly of the current state of knowledge around the topic. Five electronic databases including MEDLINE, The Cochrane Central Register of Controlled Trials, EMBASE, CINAHL and PsycINFO will be searched from inception to October 2015 (see online supplementary appendix 2 for the MEDLINE search strategy). The five databases were selected to optimise the likelihood of identifying all the previously published studies while maintaining a manageable screening load. The search uses explosions and combinations of key search terms for qualitative research, evidence-based practice, CPGs, OA, beliefs, perceptions, barriers, enablers and adherence. Search results will be collated in a reference database (Endnote VX7), duplicates deleted and then initial screening of titles will be independently conducted by two reviewers (TE and LD). A priori inclusion/exclusion criteria will be applied at this stage (see online supplementary appendix 3). The two reviewers will then independently screen abstracts of any titles retained by at least one reviewer. Inclusion/exclusion criteria will be independently applied by two reviewers to full texts of remaining references to select the final studies to include in the review. Manual searching of the reference lists and citation tracking of papers identified as potentially relevant at this stage will also be conducted. Discrepancies in the final decision on inclusion will be discussed and if necessary reviewed by a third reviewer (SCS) in order to reach consensus. Study selection will be documented and summarised in a PRISMA compliant flow chart. The process of identification and selection of articles and flow chart of article inclusion is outlined in figure 1.

Data extraction

Data extracted from the articles will include study details (location and setting, population, research question/aims, guidelines the interview questions were developed from, data collection method, data analysis method, theoretical framework) and findings (barriers, enablers, other beliefs or perspectives, second-order themes or interpretations and subthemes, supporting quotations, conclusions and recommendations; see online supplementary appendix 4A). Quotations from primary study participants (first-order data) and quotes from report authors (second-order data) will be assembled retaining links to contextual information and the findings of the assessment of quality of conduct and reporting from the original study. The extracted themes from each of the primary studies will be considered for common domains and dissonant cases. In studies where participants included patients or health professionals other than eligible primary care clinicians (eg, surgeons or rheumatologists), only data that could be attributed to the eligible primary care clinicians will be extracted. In studies that used mixed methods such as quantitative surveys, only data from qualitative components of the report will be extracted. Two reviewers (TE and LD) will independently extract the data into spreadsheets (Microsoft Excel) and discrepancies will be resolved, by consensus, when the two data sets are merged.

Critical appraisal of conduct of included studies

The Critical Appraisal Skills Programme (CASP) checklist\(^1\) will be used to identify and appraise methodological quality (trustworthiness) of the included studies independently by two raters (TE and LD). Assessment of trustworthiness in qualitative research considers study rigour, or the thoroughness and appropriateness of conduct; credibility, or whether the findings were data driven and meaningful; and relevance of the research question to the review topic.\(^43\) The CASP checklist of 10 questions comes with some decision rules and guidance on the interpretation of each item. Using the example of Slade et al,\(^44\) the review team will construct an expanded summary table detailing the presence or absence of the components of each CASP question (see online supplementary appendix 4B). A summary narrative report of the trustworthiness of the individual included studies will be provided. Summary scores or quality ratings will not be generated as the CASP does not have a scoring matrix and a cut-off point has not been established for ratings of quality of qualitative
studies. Studies will not be excluded from the review on the basis of the critical appraisal of conduct and we will discuss the impact on the data synthesis of any study weaknesses.

Data synthesis and higher order theme and theory development
A meta-synthesis approach will be used for this qualitative data synthesis, and underpinned by the philosophical positioning that knowledge of reality is mediated by one’s beliefs and perceptions. It is a systematic and comprehensive approach whereby data are coded and organised into descriptive themes, from which new higher order themes are developed that offer new interpretations beyond the primary studies’ findings.

The first stage of data synthesis will be to assemble and simplify the primary data, themes and subthemes into common groups within an external framework of barriers, enablers and other beliefs/perceptions that can act as barriers or enablers. After familiarisation with the data, recurrent codes for meaning and content across studies will be identified. Development of the codes (and a coding framework if possible) and the actual coding of the data set will be conducted by three coders (SCS, TE and LD) through discussions and iterations. All reviewers will consider all the available data. During discussions and coding rounds, ideas will be shared and refined until a final set of themes is agreed by consensus. The text to which a given code was applied will be checked for consistency of interpretation and to further refine the themes. Bearing in mind each of the broad concepts of barriers, enablers and other beliefs/perceptions relating to OA management, higher order themes will be inductively derived from the patterns in the codes. These themes may or may not have been identified by the primary study authors. Where possible, new major higher order themes will retain key contextual factors and new overarching or expanded theory will be developed through discussion between the reviewers and co-researchers. This stage relies on the judgement, insight and creativity of thinking among the researchers but will be anchored to the empirical data and review question using a grounded theory framework.

Interpretation of findings will consider new understandings in relation to factors known to influence guidelines adherence: clinician knowledge, professional background, cultural factors, environmental factors and patient drivers; and the main CPG recommendations for knee OA management related to education, self-management support for exercise and weight loss, first-line and second-line pharmacological management, and appropriate use of surgical interventions.

Assessment of credibility and rigour of new findings
The robustness of the synthesis and confidence in the review findings, defined as the analytical output of the evidence synthesis of the primary studies, will be determined as a final stage using the Confidence in the
Evidence from Reviews of Qualitative research (CERQual) approach. This approach includes consideration of (1) methodological limitations (design and conduct flaws) of each primary research study contributing to the finding and how the limitations might affect confidence in the finding; (2) relevance of the included studies to the overall review question in terms of, for example, the population, setting and original study aim; (3) coherence and consistency of results across the included studies and (4) adequacy of supporting data (quantity as well as quality). In this way, quality of the empirical studies will influence the emphasis given to data informing themes and any new theories that are generated. The CERQual is not a critical appraisal of the methodological limitations of either individual studies or the evidence synthesis, nor does it assess confidence in the overall synthesis findings, but considers each new finding separately. Confidence judgements will be achieved through discussion between at least two of the review authors. Each review finding will be allocated a level of confidence. Confidence levels start at ‘high confidence’ and are rated down by one or more levels if there are concerns regarding any of the individual CERQual components. Results will be summarised along with the main findings textually and in a table adapted from the CERQual Qualitative Evidence Profile table with the following headings:

- Individual review finding;
- Studies contributing to the review finding;
- Methodological limitations—problems with the design or conduct or reporting of primary studies;
- Relevance—applicable to context and degree to which the finding is generalisable;
- Coherence—grounding in the primary data with evidence provided by quotations from primary study participants;
- Adequacy of data—detail, depth and amount of supporting data;
- Overall CERQual assessment of confidence—four levels of confidence in the evidence for an individual review finding: high, moderate, low or very low;
- Explanation of confidence judgement.

The proposed step-by-step procedure for the data extraction and meta-synthesis is shown in figure 2.

DISCUSSION

This planned study is a systematic review and meta-synthesis that will use rigorous and explicit methods to bring together the results of empirical qualitative studies investigating perceptions and beliefs, barriers and enablers to practice based on CPG recommendations for the management of OA by primary care clinicians. The purpose is to synthesise the primary data to provide new interpretations that may assist in identification of strategies with the potential for facilitating uptake of effective treatments and CPG recommendations.
In-depth understanding of the barriers and enablers to achieving effective practice is needed in order to bridge the gap between research findings and clinical practice. Interventions which might include behaviour change interventions, service delivery changes and/or others can then be specifically designed to address the barriers unique to the population and the target practice. Implementations tailored to identify barriers may be more likely to improve practice than more general interventions. Implementing consistent and evidence-based management for patients with OA may alleviate some confusion and frustration for patients and providers, lead to better health outcomes and possibly reduce healthcare costs.

This protocol paper serves to predefine our objectives and methods and also to communicate our intent. Any deviations between this protocol and our actual methods will be discussed in the systematic review report. It is anticipated that through the assimilation and interpretation of the attitudes and experiences of primary care clinicians managing OA, reported by studies that may vary in original purpose and context, we will develop a richer understanding of the potential barriers, enablers and beliefs or perceptions that may act as barriers or enablers to optimal management of OA and assist in future policy and service delivery improvements.

Contributors TE and KB conceived the idea for the study. TE, SCS and RB were responsible for the study design and protocol. TE drafted the protocol manuscript with input from LD, SCS, RB and KB. All authors have read and approved the final manuscript. The corresponding author guarantees the paper and that the authorship statement is correct.

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REFERENCES


