


BMJ Open East Midlands knee pain multiple randomised controlled trial cohort study: cohort establishment and feasibility study protocol

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ABSTRACT

Introduction Knee pain due to osteoarthritis (OA) is a common cause of disability. The UK National Institute for Health and Care Excellence OA guidelines recommend education, exercise and weight loss advice (if overweight) as core interventions before pharmacological adjuncts. However, implementation of these in primary care is often suboptimal. This study aims to develop a complex intervention with non-pharmacological and pharmacological components that can be delivered by nurses. The feasibility and acceptability of the intervention, and feasibility of undertaking a future cohort randomised controlled trial (RCT) will be explored.

Methods and analysis In phase 1, we will develop a training programme for nurses and evaluate the fidelity and acceptability of the non-pharmacological element of the intervention. Fidelity checklists completed by the nurse will be compared with video analysis of the treatment sessions. Patients and nurses will be interviewed to determine the acceptability of the intervention and explore challenges to intervention delivery. The non-pharmacological component will be modified based on the findings. In phase 2, we will assess the feasibility of conducting a cohort RCT comprising both the pharmacological and modified non-pharmacological components. We will compare three groups: group A will receive the non-pharmacological components delivered before pharmacological components; group B will receive pharmacological components followed by the non-pharmacological components; and group C (control arm) will continue to receive usual care. Study outcomes will be collected at three time points: baseline, 13 and 26 weeks after randomisation. Qualitative interviews will be conducted with a sample of participants from each of the two active intervention arms.

Ethics and dissemination This protocol was approved by the East Midlands-Derby Research Ethics Committee (18/EM/0288) and registered at ClinicalTrials.gov (protocol v4.0, 10/02/2020). The study will be reported in accordance with the Consolidated Standards of Reporting Trials guidance and standards. The results will be submitted for publication in peer-reviewed academic journals.

Trial registration number NCT03670706

Strengths and limitations of this study

- First study to develop and evaluate the feasibility of a wholly nurse-led intervention following the core National Institute for Health and Care Excellence guidelines for treating knee pain and osteoarthritis (OA).
- This study will not determine the effectiveness of this model of care for knee pain and OA, but will explore the feasibility of implementation and running an adequately powered randomised controlled trial (RCT) and determine signal of efficacy.
- Assessing fidelity of intervention delivery will allow us to explore the extent to which the nurses can deliver individual components of the complex package of care as planned, and alongside exploring its acceptability will inform refinements to nurse training and/or the package of care.
- Blinded outcome assessment.
- Participants and nurses delivering care will not be blinded to each intervention. However, in using a cohort RCT study design where participants will be informed that their data may be used as a control comparison for intervention studies, bias associated with disappointment, behaviour modification and dropouts in the control arm will be minimised.

INTRODUCTION

Chronic knee pain affects one in four people aged 55 years or older, and is most commonly caused by osteoarthritis (OA).¹ The global prevalence of symptomatic knee OA is estimated at 4%, and it has an important impact on both affected individuals and health and social care systems.²⁻³ The socioeconomic and healthcare burden of knee OA is likely to increase due to the ageing population and the obesity epidemic.^{2,4}

Best practice guidelines for managing OA published by the National Institute for Health and Care Excellence (NICE) recommend individualised patient education, advice

and access to information about OA and its management, strengthening and aerobic exercise, and guidance on losing weight if applicable as core treatments, with consideration of adjunctive pharmacological and other non-pharmacological treatments as required.⁵ However, core treatments are frequently underused as both doctors and patients predominantly focus on pharmacological options including opioids.^{6,7} Most people with OA feel they do not receive the treatment they need from general practitioners (GP), with an overemphasis on prescription drugs and lack of discussion about exercise and diet.⁸ Given the growing concerns about the overuse of opioids for musculoskeletal pain including OA,⁹ it is important to explore alternative models of healthcare delivery with an emphasis on non-pharmacological interventions for this condition. Additionally, while it has been suggested that optimising analgesia prior to participation in exercise therapy may enhance adherence and patient outcomes, this has not been confirmed in a randomised controlled trial (RCT).¹⁰

Alternative models for implementing OA care have shown potential. The MOSAICS trial explored the effectiveness of delivering an enhanced initial consultation with GP and provision of nurse-led follow-ups as a clinically practicable way of implementing NICE guidelines compared with usual care.¹¹ However, only 29% of patients in the intervention arm reported having a consultation with a nurse, making it difficult to determine whether a nurse could help deliver the core NICE recommendations. Others have included using community physiotherapists and pharmacists for delivering interventions but have only demonstrated short-term improvements in health outcomes.¹²

The majority of patients with knee pain self-manage their symptoms, and those who seek healthcare are managed by their GPs and community physiotherapists in the first instance.⁸ A potential role for practice nurses has been identified¹³ and nurse-led clinics already exist for patients with long-term conditions such as coronary heart disease,¹⁴ heart failure¹⁵ and diabetes¹⁶ resulting in equivalent or better outcomes for patients compared with usual GP-led care. Upskilling practice nurses to deliver the management of long-term conditions is recognised as a key strategy for the future of primary care.¹⁷ This paper describes the protocol for developing and testing the feasibility of a wholly nurse-led intervention for people with knee pain, delivering the core NICE recommendations.

Further to this, it will test the feasibility of a cohort RCT. The usual double-blind, placebo-controlled RCT cannot be used when investigating the efficacy of complex interventions for knee pain as it is not possible to blind participants to their treatment. This has led to the use of pragmatic RCTs where participants are randomised to receive an intervention or continue usual primary care.^{12,18} Such studies can be affected by disappointment bias, behaviour modification bias and differential dropouts when those selected as controls had hoped to receive the intervention under investigation and drop out

when allocated to the control arm. Cohort RCTs, as will be used in this study, have been adopted to prevent such biases.^{19,20} In this study design, a cohort of people with the condition of interest are recruited and consented to be approached to complete further questionnaire surveys, for their GP medical records and prescriptions to be accessed, to be used as a comparator for future studies and to be approached for participation in future studies if eligible.²¹ Clinical and patient-reported outcomes are captured at regular intervals, allowing those eligible for any given study to act as a control without being informed about the experimental intervention and thus minimising the bias associated with pragmatic trials.

Aims and objectives

The overall purpose is to develop and test the feasibility of a nurse-led intervention for people with knee pain using a cohort RCT study design.

This study has two phases:

Phase 1 involves the development and evaluation of the non-pharmacological treatment component. Specific objectives of phase 1 are to: (1) develop a training package for nurses to deliver the core non-pharmacological and pharmacological principles to manage knee OA as recommended by NICE; (2) determine the fidelity of delivery of the nurse-delivered components of the intervention; (3) explore patient and nurse acceptability of the non-pharmacological components of the intervention.

Phase 2 will test the feasibility of a cohort RCT of nurse-led versus usual care of people with knee pain and explore whether such a trial should provide analgesia before non-pharmacological interventions.

METHODS

Participants

Participants eligible for both phases of the study will be aged 40 years or older and self-reporting knee pain on most days of the previous month for at least 3 months. Knee pain severity will be scored between 4 and 7 on a 0–10 numeric rating scale (NRS). This will be assessed using the following question: ‘Over the past 4-weeks, how intense was the average pain or aching in your knees on a 0–10 scale, where 0 is no pain and 10 is pain as bad as could be?’

Exclusion criteria include participants who are unable to communicate in English, who are housebound or care home residents, on dialysis or home oxygen, pregnant or have dementia, serious mental illness, terminal cancer, autoimmune rheumatic diseases, asthma or lung disease requiring regular daily oral corticosteroids, unstable angina or heart failure, known peripheral vascular disease, stroke with residual weakness or sensory loss, physician-diagnosed peripheral neuropathy with sensory or motor deficit, previous knee or hip replacement, on a waiting list for a knee or hip replacement or with severe knee pain (score >7 on NRS for average pain in the past 4 weeks).

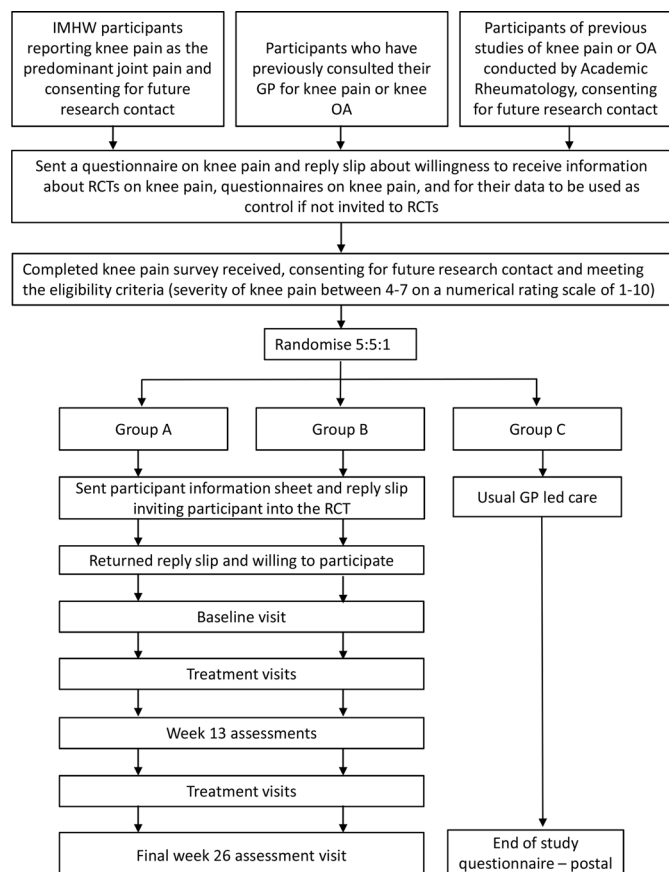


Figure 1 Participant timeline through the study. GP, general practitioner; IMHW, Investigating Musculoskeletal Health and Wellbeing cohort study; OA, osteoarthritis; RCT, randomised controlled trial.

Nurses who undergo the intervention training and delivered the intervention in the study will be recruited to the qualitative components of the study and will give their consent to be interviewed in both phases of the study.

Recruitment

There are three routes of recruitment:

1. Participants will primarily be recruited from the Investigating Musculoskeletal Health and Wellbeing survey (NCT03696134), a cohort study that aims to measure and characterise the development and progression of pain, frailty and disability and form a longitudinal context for nested research.²¹ Participants who self-report knee pain as their predominant body pain and consent for future research contact will be sent a questionnaire enquiring about their knee pain, mood, function and quality of life. They will also be asked about willingness to receive information about trials on knee pain, complete further questionnaires on knee pain, and for their data to be used for comparisons with other participant groups in research studies. Those willing and meeting the eligibility criteria will be invited to take part in the current study (figure 1).
2. Additional participants identified by screening of GP records for previous consultation for knee pain will be sent a questionnaire as outlined above.

3. Finally, people with knee pain or OA who have taken part in previous community-based surveys in Academic Rheumatology, University of Nottingham and consented for future research contact may be approached with a questionnaire.

Setting

Study visits for research assessment will occur at Academic Rheumatology, City Hospital Nottingham, UK and the David Greenfield Unit, Queen's Medical Centre, Nottingham, UK. The intervention will be delivered at Academic Rheumatology, City Hospital Nottingham.

Consent and withdrawal

All participants will give written informed consent before entering the study and before any assessments or interventions related to the study are undertaken. Optional consent will be sought for video recording of the intervention sessions to evaluate fidelity and audio recording of participant interviews. Participants will be free to withdraw at any time if they desire to do so, or at the discretion of the chief investigator. In the event of withdrawal, any data collected up until that point will be kept and potentially included in any analyses.

Patient and public involvement

The design of this study was supported by a patient advisory group of three people with hip or knee OA and a range of experiences in primary and secondary care. The group recognised the challenges of providing full explanations and individualised advice in a time-limited GP consultation and agreed that a wholly nurse-led intervention would be acceptable or preferable to most people. They provided input into the content of the both pharmacological and non-pharmacological components, the number and length of intervention sessions and the use of exercise diaries and educational content.

PHASE 1: DEVELOPMENT OF THE NON-PHARMACOLOGICAL COMPONENT

The non-pharmacological intervention will be developed according to the Medical Research Council framework for developing complex packages of care.²² It was informed by several strands of evidence including current guidelines, an expert multidisciplinary team of physiotherapists, rheumatologists and nurses, patient opinion, physiological considerations and behaviour change theory.

In brief, the intervention consists of a holistic assessment of the participant, individualised education about OA, aerobic and strengthening exercises and weight loss advice if required. Evidence-based strategies to motivate participants and support adherence to healthy behaviours will be employed.

The training package

A training package for nurses will be developed and piloted during this phase. Research nurses will be recruited

without any prior knowledge of treating musculoskeletal conditions. The content will be based on current NICE guidelines for the management of OA⁵ and a report by Arthritis Research UK on the educational needs of health professionals working with people with OA.²³ Academic and clinical experts and members of a patient and public involvement group have provided input into the training package.

Delivery of the training package will be underpinned by educational theory²⁴ and will build on the nurses existing knowledge of the condition. Training will be delivered in face-to-face sessions, complemented by a range of learning resources including a manual, case studies, online resources and patient simulations. Key components of the training will include:

- ▶ The epidemiology and nature of knee pain and knee OA.
- ▶ Assessment of the patient with knee OA.
- ▶ Core NICE guidelines for managing OA.
- ▶ Principles of strengthening and aerobic exercise prescription for knee OA.
- ▶ Information and advice to support weight loss.
- ▶ Strategies to support behaviour change.
- ▶ Pharmacological management of OA and knee pain following a stepwise protocol of optimising analgesia.

Study design

A pre-post intervention study using mixed quantitative and qualitative methods to determine the fidelity of delivery and the acceptability of the intervention by patients and nurses. After the nurse training, 20 participants with knee pain will be recruited to this study. They will receive the non-pharmacological components of the intervention covering as described above in four sessions over a 5-week period.

Quantitative methods

Fidelity of delivery refers to the assessment of whether content of the sessions was delivered as intended. This will be assessed using a priori fidelity checklist of five to eight categories and individual components (online supplementary additional file 1). Categories included materials, introduction, assessment, education, exercise, weight loss, adjunct treatments, and review and planning. All interventions in this phase will be video recorded (with consent). Nurses will self-complete the checklist after each session and a researcher will independently complete the checklist using the video recordings. The individual components of each category for each session will be rated as being complete, partially complete, not completed or not applicable. A sample of video recordings will be reviewed by a second researcher to determine inter-rater reliability.

Quantitative analysis

Fidelity scores will be presented as the percentage of components that were delivered as intended for the overall delivery of the intervention, for each session and

for each category. Inter-rater reliability between the two researchers scoring the video recordings will be reported, as will the level of agreement between nurse-completed scores and researcher-completed (video) scores. For delivery of complex interventions such as this, levels of fidelity have been previously interpreted as ‘high’ fidelity where 80%–100% of the specified components were delivered as intended, ‘moderate’ 51%–79% and ‘low’ 0%–50%.²⁵ Where the fidelity scores are less than 80%, we will explore further to establish which components are responsible.

Qualitative methods

Acceptability of the intervention will be explored in a face-to-face interview with all participants who received the intervention. Participants who withdraw from the intervention will be offered the opportunity to take part in an interview to explore their experiences and reasons for discontinuation. The nurses who deliver the intervention will also be invited to interview, to explore their views on the training, experience in delivering the intervention and perceived factors influencing the fidelity of delivery. Interviews will be carried out by a PhD student (PAN) and overseen by two experienced qualitative researchers (AF and RdN). All interviews will be audio recorded and transcribed verbatim by an external specialist company Transcribe It and anonymised.

Qualitative analysis

After data transcription, all data will be checked for accuracy before transcripts are imported to NVivo V.12. Qualitative data will be analysed using a framework approach.²⁶ This method sits within the broad family of thematic analysis, but is particularly useful for research that has specific questions and a priori issues that need to be dealt with.²⁷ The analysis will follow the five stages of framework analysis: familiarisation with the data, construction of an initial thematic framework, indexing and sorting the data using initial thematic framework, finalisation of thematic framework, and summarising and displaying the data into a matrix. Emergent themes and subthemes will be discussed and agreed by at least two researchers to increase the validity of the analyses.

Following the fidelity evaluation and qualitative interviews, modifications may be made to study materials, procedures or protocol, and/or nurse training.

PHASE 2: FEASIBILITY COHORT RCT

Trial design

This will be a single-centre, mixed methods feasibility cohort RCT. Participants will be recruited as described above and randomised to one of three treatment arms (figure 1).

Group A will receive the non-pharmacological protocol for 13 weeks followed by the pharmacological protocol between weeks 13 and 26 as required.

Group B will receive the pharmacological protocol in the first 13 weeks followed by the non-pharmacological protocol between weeks 13 and 26 with optimised background analgesia.

Group C is a control (cohort) group and will continue to receive usual care.

Randomisation and allocation concealment

Participants will be individually randomised on a 5:5:1 ratio using randomly permuted block sizes of 3 and 6, stratified for the number of eligible knees (ie, unilateral or bilateral knee pain). Randomisation codes will be generated by the study statistician. Allocations to groups will be enclosed in serially numbered, opaque, sealed envelopes with a carbon copy paper. The serially numbered opaque envelopes will be packaged and prepared by an independent member not belonging to the research team. Participants will be randomised by the trial coordinator who will ensure that the envelopes are opened sequentially, and only after the participant's unique study identifier is written on the outside of the appropriate envelope.

Blinding

It is not possible to blind the study participants or the nurse delivering the intervention to the group allocations. However, study personnel involved in outcome assessment and data analysis will be blinded. Participants will be requested not to disclose group allocation to the outcome assessor, but if this does occur it will be recorded. Only once data have been cleaned and analysed will the treatment allocation be made known.

Intervention

The intervention will be delivered by the same nurse as in phase 1, any additional nurses recruited will undergo the same training.

Non-pharmacological component

The non-pharmacological component will be delivered by a nurse as detailed in [table 1](#) incorporating any modifications made following the development phase (a detailed description is included in online supplementary additional file 2). It will be delivered in up to six face-to-face sessions over 13 weeks. Participants will continue their usual analgesics in this period.

Pharmacological component

The nurse will take a history of current symptoms, comorbidities, medications and the main knee complaint of the participant. They will be advised by the nurse to continue on their current analgesia prescribed by their GP, and to add in simpler and safer analgesics in the sequence shown in [figure 2](#). Once they are on a simpler and safer analgesic, they will be advised to reduce the dose of potentially more toxic or stronger analgesic. However, this will depend on the subjective improvement the participant experiences with simpler analgesic.

Relative and absolute contraindications will be assessed against a checklist developed from the British National

Formulary. Participants with absolute contraindication to an analgesic will not be prescribed that drug. Participants with relative contraindications may be prescribed the aforementioned drug if they are willing to do so after exploring risks and benefits.

Analgesics will be reviewed at two weekly intervals over a 13-week period and optimised if the pain relief is insufficient. This will be done over the phone, or at a face-to-face visit, depending on participant preference. The visits and telephone consultations will be conducted by the nurse and prescriptions signed by the principal investigator or nominated deputy. Once a participant achieves adequate pain control and does not request any further changes to their analgesia, they will be advised to contact the nurse by phone for changes to their treatment if needed during the study.

Usual care

Participants allocated to this group will continue to receive usual care for their knee pain. They will not undertake any of the clinical assessments and will not receive any input from the nurse. People in this group will be part of the baseline cohort and will not be aware of the content of the intervention groups.

Concomitant treatments

Participation in the trial does not preclude the participants from receiving any concomitant care or treatment.

Quantitative study outcomes

The feasibility of running a full trial will be assessed by recording the following data:

- ▶ Recruitment rates.
- ▶ Dropout rate and reasons for dropout.
- ▶ Number of scheduled nurse appointments attended.
- ▶ Number of instances of unblinding.
- ▶ Completeness of questionnaire data.
- ▶ Concordance with exercise assessed using data from participants' exercise diaries (total number of days on which exercises were performed).

Participant-reported measures

A summary of all participant outcomes to be collected at 0, 13 and 26 weeks for groups A and B is presented in [table 2](#) and in online supplementary additional file 3.

Group C will receive a questionnaire at week 26, enquiring about healthcare utilisation, demographic characteristics, self-reported height, self-reported weight, current alcohol intake and smoking status, comorbidities, medications, joint pain, central aspects of pain in knee scale, Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC), Short Form-36 Health Survey version 2 and Hospital Anxiety and Depression Scale.

Safety and adverse events

This study intervention follows the current NICE guidelines that might be offered as part of routine clinical care. As such the risk of severe or unexpected adverse

Table 1 Content of non-pharmacological component of intervention

Content of non-pharmacological component	Session					
	1	2	3	4	5	6
Assessment						
Holistic assessment including symptoms, pain elsewhere, comorbidities, impact on function, occupation, mood, sleep, illness perceptions, current levels of PA and attitudes to PA and weight loss (if required)	✓					
Physical assessment of knee range of movement, lower limb muscle strength, observation of gait and functional activities, BMI	✓					
Education and advice						
Provision of Arthritis Research UK booklet 'Osteoarthritis of the knee'	✓					
Nature of osteoarthritis	✓					
Adverse illness perceptions addressed	✓					
Core treatments	✓					
Benefits of exercise and PA	✓					
Pacing	✓					
Benefits of weight loss (if required)	✓					
Use of heat and cold for pain	✓					
Appropriate footwear, use of walking aids	✓					
Signposting to further information if required	✓					
Review of above if required		✓	✓	✓	✓	✓
Exercise (individualised programme)						
Strengthening exercise	✓					
Aerobic exercise/PA	✓					
Functional exercises	✓					
Stretching exercises	✓					
Review performance of exercise		✓	✓	✓	✓	✓
Progression/regression of exercises		✓	✓	✓	✓	✓
Weight loss (if required)						
Previous efforts to lose weight discussed	✓					
Strategies for weight loss discussed	✓					
Agree weight loss goal (5% body weight)	✓					
Signposting to resources (weight loss groups, NHS weight loss plan)	✓					
Review of weight loss progress and advice		✓	✓	✓	✓	✓
Adherence/behaviour change strategies						
Patient goals and action plan recorded for exercise and weight loss	✓	✓	✓	✓	✓	✓
Patient's confidence to achieve goals discussed and recorded	✓	✓	✓	✓	✓	✓
Barriers and facilitators discussed	✓	✓	✓	✓	✓	✓
Exercise diary completed and discussed	✓	✓	✓	✓	✓	✓

BMI, body mass index; NHS, National Health Service; PA, physical activity.

events (AE) is low.²⁸ Exercise and an increase in physical activity may initially increase the risk of AEs such as pain, fatigue or muscle soreness or increased falls through increased activity. To reduce the risk of AEs the exercise programme will be tailored to the abilities of the participants. All serious and non-serious AEs will be monitored and recorded through the study by

the nurses and will be managed in line with current National Institute for Health Research guidelines.²⁹

Qualitative study outcomes

Acceptability of the intervention will be assessed using qualitative interviews conducted after the intervention (week 26) with approximately 10 participants

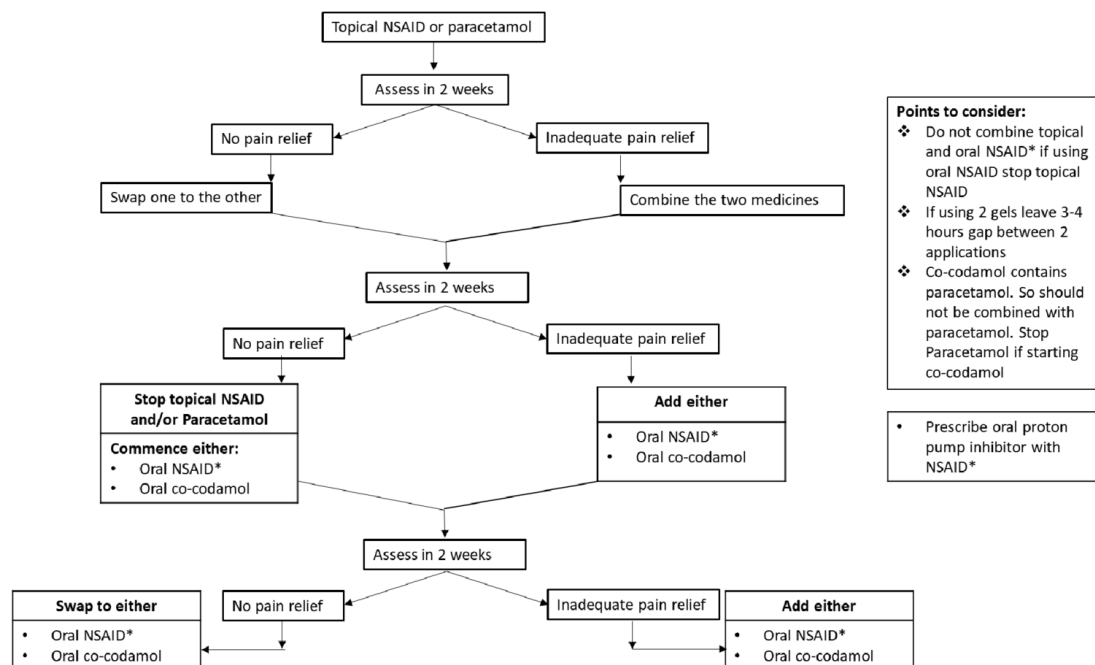


Figure 2 Analgesic sequence for pharmacological component. NSAID, non-steroidal anti-inflammatory drug.

from each of the intervention arms (groups A and B). They will be purposively selected to represent those with likely low and high concordance with the exercise advice using the Adherence to Exercise Scale for Older Patients questionnaire. Interviews will explore participants' overall satisfaction with the intervention and the sequence of treatment, perceptions of nurse-led care and previous treatment experience, level of adherence to the advice, perceptions of managing their knee pain, as well as perceived impact of their knee pain on their daily life before and after the intervention. Participants who withdraw from the intervention will be offered the opportunity to take part in an interview. Interviews will also be conducted with the study nurses to explore their experience in delivering the intervention, perceived effectiveness of the intervention and barriers to implementation and how these may be overcome.

Sample size

Quantitative study

As this is a feasibility study, a formal sample size calculation for between-group comparisons of primary clinical outcome is not appropriate. A target sample size of 53 participants per arm will be sought over the recruitment period to reliably estimate the feasibility outcomes relating to recruitment and retention rates to inform a fully powered RCT. With a sample size of 159 (53 participants per arm), we will be able to estimate a dropout rate of no more than 20% to within 7% points of the true value with 95% confidence.

Qualitative study

Target recruitment will be 20 participants, 10 from each intervention arm. However, final numbers will be

determined by data saturation, where no new themes are identified.

Data analysis

Feasibility outcomes will be estimated using descriptive statistics (with 95% CIs) and will be presented overall and per randomised groups. A Consolidated Standards of Reporting Trials (CONSORT) diagram will summarise the flow of participants through the study. Reasons for non-eligibility, withdrawals and non-completion of follow-up questionnaires will be presented if available. As this is a feasibility study, interim analyses are not planned. Similarly, missing data will not be imputed; on the contrary, the pattern of missing data will be assessed. Protocol non-adherence will be assessed as randomised.

Quantitative analysis

The main quantitative analysis will be of the trial feasibility outcomes:

- ▶ Recruitment rate (per month, per recruitment source, per 100 participants approached).
- ▶ Dropout rate (per arm, per stage pharmacological/non-pharmacological).
- ▶ Attendance rates for scheduled nurse appointments.
- ▶ Missing data.
- ▶ Power and sample size calculation for a definitive trial will be based on WOMAC summated knee pain domain scores in the most painful knee at week 26.

Descriptive statistics will be presented for demographic data and all baseline clinical outcome measures. Exploratory analysis of clinical outcomes will be conducted according to randomised groups but will not be interpreted in terms of effectiveness. The emphasis will be on CIs of effect size estimations rather than the p values. Changes

**Table 2** Summary of questionnaire and research measures to be collected

Domain	Data source	Measure/instrument	Time points (week)
Demographic characteristics	Research assessment	Age, height, weight	0 0,13,26
Comorbidities and medications	Research assessment	Comorbidities, current medications	0 0,13,26
Radiographic evaluation	Research assessment	Bilateral knee radiographs: PA semiflexed weight bearing and skyline views ³⁷ scored for Kellgren and Lawrence grades and using the Nottingham line drawing atlas ³⁸	0
Knee pain	Self-report	Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC) ³⁹	0,13,26
	Self-report	Analgesic and NSAID consumption	0,13,26
	Research assessment	Quantitative sensory testing* including pressure pain detection threshold, temporal summation and conditioned pain modulation	0,13,26
Physical activity	Self-report	International Physical Activity Questionnaire (IPAQ) ⁴⁰	0,13,26
Function	Self-report	WOMAC function subscale ³⁹	0,13,26
	Research assessment	Timed Up and Go (TUG) test ⁴¹ 30s chair stand test ⁴²	0,13,26
Muscle function	Research assessment	Isometric and isokinetic quadriceps strength*	0,13,26
Quality of life	Self-report	The Short Form-36 Health Survey V2 ⁴³ EQ-5D-5L ⁴⁴	0,13, 26
Psychological wellness	Self-report	Hospital Anxiety and Depression Scale (HADS) ⁴⁵	0,13,26
Healthcare use	Self-report	Service use questionnaire to assess use of NHS or private healthcare, prescription and over-the-counter medicines related to knee pain outside of the study	0,13,26
Participant engagement in treatment	Nurse-completed questionnaire	Pittsburgh Rehabilitation Participation Scale (PRPS) ⁴⁶	13/26†
Exercise adherence	Self-report	Adherence to Exercise Scale for Older Patients (AESOP) ⁴⁷	13/26‡
Acceptability of intervention	Self-report	Participant satisfaction with treatment	13,26
Blood markers	Research assessment	Non-fasting serum cholesterol, HbA1c and C-reactive protein§	0,13,26
Safety	Case report form	Adverse events	26

*Further description included in detail in online supplementary additional files 3 and 4.

†This will be completed by the nurse after each of the non-pharmacological sessions.

‡This will be completed by the participant at the end of non-pharmacological arm.

§Online supplementary additional file 4.

EQ-5D-5L, 5-level version of EuroQol-5 Dimension; NHS, National Health Service; NSAID, non-steroidal anti-inflammatory drug; PA, physical activity.

in clinical and patient-reported outcomes from baseline to 13 and 26 weeks will be analysed using appropriate parametric or non-parametric statistics. A comparison of those receiving the pharmacological component first with those receiving the non-pharmacological component first will help determine the order of delivery in a future trial.

Qualitative analysis

Interview data will be analysed following the framework approach as described in phase 1.³⁰ Analysis will be conducted in parallel with the interviews and initial results will inform subsequent sampling and areas of interest to follow-up.

CRITERIA FOR TERMINATING THE STUDY

The study may be stopped by the sponsor if there is apparent futility in continuing with it.

DATA MANAGEMENT

Study data will be managed by the study coordinator (BM) under the supervision of the chief investigator (AA) and the study statistician (RO). A data monitoring committee has not been convened by the sponsor as this is a feasibility study using well-established and NICE-approved interventions. All trial staff and investigators will endeavour to protect the rights of the trial's participants to privacy and informed consent, and will adhere to the Data Protection Act 2018 and the General Data Protection Regulation (European Union) 2016/679. The study coordinator shall carry out monitoring of trial data as an ongoing activity. Trial data and evidence of monitoring and systems audit will be made available for inspection as required by the sponsor and the Research Ethics Committee (REC).

Qualitative data and video or audio recordings will be archived in the University of Nottingham servers using password protection. Quantitative trial data will be stored in Research Electronic Data Capture database with range checks for data values. All quantitative data will be source data verified. Ten per cent of the entered trial data will be audited and variables for which there is >5% error will be entered again. Once auditing is complete the electronic case report forms will be signed off and the database placed under hard lock. Study data will be available to the study statistician, PhD student and research fellow working on the project.

In compliance with the International Conference on Harmonization/Good Clinical Practice guidelines, regulations and in accordance with the University of Nottingham Research Code of Conduct and Research Ethics, the chief or local principal investigator will maintain all records and documents regarding the conduct of the study. These will be retained for at least 7 years or longer if required.

ROLE OF SPONSOR AND FUNDERS

There is no role of the study sponsors and funders in the design; collection, management, analysis and interpretation of data; writing of the report; and the decision to submit the report for publication. The study sponsor and funder will have no ultimate authority over any of these activities.

ETHICS AND DISSEMINATION

This protocol was given approval approved by the East Midlands-Derby REC (18/EM/0288) prior to commencing recruitment in November 2018 which is ongoing. The study was registered at ClinicalTrials.gov. The trial will be reported according to CONSORT guidance and standards.³¹ The results will be submitted for publication in peer-reviewed academic journals. Any modification to the approved protocol will result in resubmission to gain approval from the REC and study sponsor. Authorship eligibility guidelines will be adhered to, and we do not intend to use professional writers. Any interested researcher will be able to access the full protocol, participant-level data set and statistical code for research purposes after data sharing agreement has been signed.

DISCUSSION

People with knee OA continue to often receive suboptimal fragmented care and the core NICE recommendations are underused in primary care.³² Given that knee pain is common, and there is a huge time pressure on GPs in the UK, it is vital to find out if a complex package of care incorporating the core recommendations can be delivered by other healthcare professionals such as practice nurses.

We believe that a wholly nurse-led management programme where a nurse acts as the point of contact for people with knee pain due to OA, educates them about the condition, provides core pharmacological and non-pharmacological treatments and builds a long-term therapeutic relationship is likely to be

clinically effective and cost-effective in improving quality of OA care as demonstrated for other chronic conditions.^{16 33–36}

This study will not provide an answer as to the effectiveness of a wholly nurse-led intervention for OA and knee pain, but will determine the feasibility of implementing this model of care and of testing it in a full trial using practice nurses. Assessing fidelity will play an important part in the development and evaluation of the intervention, by exploring the extent to which the nurses can deliver individual components of the complex package of care for knee pain as planned. Combined with qualitative interviews to explore their acceptability, this will provide insight into the suitability of the training and intervention to inform appropriate refinements to the whole package of care that can be tested in a full trial. Although the intervention will take place in a research setting rather than primary care, the study outcomes will provide insight into the feasibility of implementation into real-world practice.

Further, this study will explore at what point analgesia should be optimised within this complex package of care. Having two intervention groups, one where analgesia is provided before the non-pharmacological component and one where it is provided after, will help us determine whether patients exercise better when analgesia is optimised first, or whether they are able to exercise sufficiently before this. This will inform the order of treatment in a two-arm full trial.

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