Improving the self management of chronic pain

Submission date Recruitment status [X] Prospectively registered

06/04/2011 No longer recruiting [X] Protocol

Registration date Overall study status [X] Statistical analysis plan

05/05/2011 Completed [X] Results

Last Edited Condition category Individual participant data

11/06/2018 Musculoskeletal Diseases

Plain English summary of protocol

Not provided at time of registration

Study website

http://www.icms.qmul.ac.uk/chs/pctu/current_projects/copers

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

V10 10.1.11, 69838

Study information

Scientific Title

COping with persistent Pain, Effectiveness Research in Self-management

Acronym

COPERS

Study objectives

Null Hypothesis - There is no difference in pain related disability between those exposed to the self-management chronic pain course and those with usual GP care plus relaxation.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Cambridgeshire Research Ethics Committee 4, 18/03/2011, ref: 11/EE/0046

Study design

Pragmatic randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

GP practice

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use contact details to request a participant information sheet

Health condition(s) or problem(s) studied

Chronic musculoskeletal pain

Interventions

- 1. To test the effectiveness of a self-management course for chronic pain against a control consisting of usual GP care, a patient education leaflet and a relaxation CD.
- 2. The intervention is a group based, facilitated learning course about coping strategies for living with chronic pain
- 3. The course is led by a health care professional and a lay person with chronic pain. We aim to have around 12 participants per course
- 4. The course will cover various aspects of pain education, pain management techniques, posture

and movement

5. It will be run over three days within one week with a two hour follow up session after two weeks

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Pain-related disability

Secondary outcome measures

- 1. Health economics: Incremental Cost Utility Ratio (ICUR)
- 2. Coping skills, anxiety, depression, social integration and self efficacy

Overall study start date

01/06/2011

Completion date

23/07/2012

Eligibility

Key inclusion criteria

Adults (aged 18 or over) with chronic musculoskeletal pain

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

700 (703 participants by end of recruitment)

Key exclusion criteria

- 1. Not fluent in English
- 2. Serious active co-morbidity that is more disabling to the individual than chronic pain
- 3. Serious mental health issues that would make it difficult for an individual to participate in the group course
- 4. Patients with a life expectancy of less than six months
- 5. Substance misuse that would make it difficult for an individual to participate in the group

course
6. Inability to give informed consent

Date of first enrolment

01/06/2011

Date of final enrolment

23/07/2012

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
Queen Mary University of London
London

United Kingdom E1 2AT

Sponsor information

Organisation

Queen Mary University of London (UK)

Sponsor details

Joint Research and Development Office Queen Mary Innovation Centre 5 Walden Street London England United Kingdom E1 2EF

Sponsor type

University/education

ROR

https://ror.org/026zzn846

Funder(s)

Funder type

Government

Funder Name

National Institute for Health Research (NIHR) (UK) Programme Grants for Applied Research (RP-PG-0707-10189)

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Protocol article	protocol	28/01/2013		Yes	No
Results article	results	14/11/2013		Yes	No
Results article	results	15/11/2013		Yes	No
Statistical Analysis Plan	statistical analysis plan	15/02/2014		No	No
Results article	results	14/06/2016		Yes	No
Results article	cohort analysis results	06/06/2018		Yes	No