Randomised single blind controlled trial to evaluate the use of self timed paced functional activity in chronic low back pain patients

Submission date 30/09/2005	Recruitment status No longer recruiting	 Prospectively registered Protocol
Registration date 30/09/2005	Overall study status Completed	 Statistical analysis plan Results
Last Edited 26/03/2020	Condition category Musculoskeletal Diseases	 Individual participant data Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s) Scientific

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers N0241139833

Study information

Scientific Title

Randomised single blind controlled trial to evaluate the use of self timed paced functional activity in chronic low back pain patients

Study objectives

Does the use of a stopwatch in timing paced physical function significantly reduce self-reports of disability and pain in the chronic low pain population

Ethics approval required Old ethics approval format

Ethics approval(s) Not provided at time of registration

Study design Randomised controlled trial

Primary study design Interventional

Secondary study design Randomised controlled trial

Study setting(s) Not specified

Study type(s) Not Specified

Participant information sheet

Health condition(s) or problem(s) studied Musculoskeletal Diseases: Low back pain

Interventions

40 chronic low back pain patients will be selected using the inclusion and exclusion criteria from the physiotherapy waiting list in the Pain Clinic at St Marys

Intervention Type Other

Phase Not Specified

Primary outcome measure

The following outcome measures will be utilised by using Short Form Mc Gill Pain Questionnaire, Oswestry Disability Questionnaire and Tampa Scale for Kinesiophobia Questionnaire **Secondary outcome measures** Not provided at time of registration

Overall study start date 30/11/2003

Completion date 30/04/2004

Eligibility

Key inclusion criteria 40 patients

Participant type(s) Patient

Age group Not Specified

Sex Not Specified

Target number of participants 40

Key exclusion criteria Does not match inclusion criteria

Date of first enrolment 30/11/2003

Date of final enrolment 30/04/2004

Locations

Countries of recruitment England

United Kingdom

Study participating centre St Mary's NHS Trust London United Kingdom W2 1NY

Sponsor information

Organisation Department of Health

Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL +44 (0)20 7307 2622 dhmail@doh.gsi.org.uk

Sponsor type Government

Website http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type Hospital/treatment centre

Funder Name St Mary's NHS Trust (UK) NHS R&D Support Funding

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